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NEW BISPIDINE COMPOUNDS USEFUL IN THE TREATMENT OF  
CARDIAC ARRHYTHMIAS

**Field of the Invention**

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This invention relates to novel pharmaceutically useful compounds, in particular compounds which are useful in the treatment of cardiac arrhythmias.

10 **Background and Prior Art**

Cardiac arrhythmias may be defined as abnormalities in the rate, regularity, or site of origin of the cardiac impulse or as disturbances in conduction which causes an abnormal sequence of activation. Arrhythmias may be  
15 classified clinically by means of the presumed site of origin (i.e. as supraventricular, including atrial and atrioventricular, arrhythmias and ventricular arrhythmias) and/or by means of rate (i.e. bradyarrhythmias (slow) and tachyarrhythmias (fast)).

20 In the treatment of cardiac arrhythmias, the negative outcome in clinical trials (see, for example, the outcome of the Cardiac Arrhythmia Suppression Trial (CAST) reported in New England Journal of Medicine, 321, 406 (1989)) with "traditional" antiarrhythmic drugs, which act primarily by slowing the conduction velocity (class I antiarrhythmic drugs), has prompted  
25 drug development towards compounds which selectively delay cardiac repolarization, thus prolonging the QT interval. Class III antiarrhythmic drugs may be defined as drugs which prolong the trans-membrane action potential duration (which can be caused by a block of outward K<sup>+</sup> currents

or from an increase of inward ion currents) and refractoriness, without affecting cardiac conduction.

One of the key disadvantages of hitherto known drugs which act by delaying  
 5 repolarization (class III or otherwise) is that they all are known to exhibit a  
 unique form of proarrhythmia known as *torsades de pointes* (turning of  
 points), which may, on occasion be fatal. From the point of view of safety,  
 the minimisation of this phenomenon (which has also been shown to be  
 exhibited as a result of administration of non-cardiac drugs such as  
 10 phenothiazines, tricyclic antidepressants, antihistamines and antibiotics) is a  
 key problem to be solved in the provision of effective antiarrhythmic drugs.

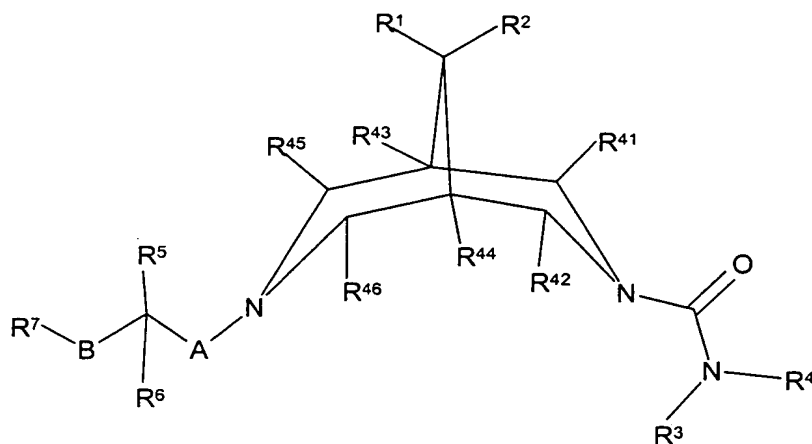
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 Antiarrhythmic drugs based on bispidines (3,7-diazabicyclo[3.3.1]nonanes),  
 are known from *inter alia* international patent application WO 91/07405,  
 15 European patent applications 306 871, 308 843 and 655 228 and US patents  
 3,962,449, 4,556,662, 4,550,112, 4,459,301 and 5,468,858, as well as  
 journal articles including *inter alia* J. Med. Chem. **39**, 2559, (1996),  
 Pharmacol. Res., **24**, 149 (1991), Circulation, **90**, 2032 (1994) and Anal.  
 Sci. **9**, 429, (1993). Known bispidine-based antiarrhythmic compounds  
 20 include bisaramil (3-methyl-7-ethyl-9 $\alpha$ ,4'-(Cl-benzoyloxy)-3,7-  
 diazabicyclo[3.3.1]nonane), tedisamil (3',7'-bis(cyclopropylmethyl)spiro-  
 (cyclopentane-1,9')-3,7-diazabicyclo[3.3.1]nonane), SAZ-VII-22 (3-(4-  
 chlorobenzoyl)-7-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane), SAZ-VII-23  
 (3-benzoyl-7-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane), GLG-V-13 (3-[4-  
 25 (1H-imidazol-1-yl)benzoyl]-7-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane),  
 KMC-IV-84 (7-[4'-(1H-imidazolo-1-yl)benzenesulfonyl]-3-*iso*-propyl-3,7-  
 diazabicyclo[3.3.1]nonane dihydroperchlorate and ambasilide (3-(4-  
 aminobenzoyl)-7-benzyl-3,7-diazabicyclo[3.3.1]nonane).

We have surprisingly found that a novel group of bispidine-based compounds exhibit electrophysiological activity, preferably class III electrophysiological activity, and are therefore expected to be useful in the treatment of cardiac arrhythmias.

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## Disclosure of the Invention

According to the invention there is provided compounds of formula I,



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wherein

$R^1$  and  $R^2$  independently represent H,  $C_{1-4}$  alkyl,  $OR^{2b}$  or  $N(R^{2c})R^{2d}$ , or  
15 together form  $-O-(CH_2)_2-O-$ ,  $-(CH_2)_3-$ ,  $-(CH_2)_4-$  or  $-(CH_2)_5-$ ;

$R^{2b}$ ,  $R^{2c}$  and  $R^{2d}$  independently represent H or  $C_{1-6}$  alkyl;

$R^3$  represents H,  $C_{1-6}$  alkyl or, together with  $R^4$ , represents  $C_{3-6}$  alkylene  
(which alkylene group is optionally interrupted by an O atom and/or is  
20 optionally substituted by one or more  $C_{1-3}$  alkyl groups);

$R^4$  represents H,  $C_{1-12}$  alkyl,  $C_{1-6}$  alkoxy (which latter two groups are both  
optionally substituted and/or terminated by one or more substituents  
selected from -OH, halo, cyano, nitro,  $C_{1-4}$  alkyl and/or  $C_{1-4}$  alkoxy),

$-(CH_2)_q$ -aryl,  $-(CH_2)_q$ -oxyaryl,  $-(CH_2)_q$ -Het<sup>1</sup> (which latter three groups are optionally substituted (at the  $-(CH_2)_q$ - part and/or the aryl/Het<sup>1</sup> part) by one or more substituents selected from -OH, halo, cyano, nitro,  $-C(O)R^{10}$ ,  $-C(O)OR^{11}$ ,  $-N(H)S(O)_2R^{11a}$ ,  $C_{1-6}$  alkyl and/or  $C_{1-6}$  alkoxy),

- 5  $-(CH_2)_qN(H)C(O)R^8$ ,  $-(CH_2)_qS(O)_2R^8$ ,  $-(CH_2)_qC(O)R^8$ ,  $-(CH_2)_qC(O)OR^8$ ,  $-(CH_2)_qC(O)N(R^9)R^8$  or, together with  $R^3$ , represents  $C_{3-6}$  alkylene (which alkylene group is optionally interrupted by an O atom and/or is optionally substituted by one or more  $C_{1-3}$  alkyl groups);

q represents 0, 1, 2, 3, 4, 5 or 6;

- 10  $R^8$  represents H,  $C_{1-6}$  alkyl, aryl (which latter group is optionally substituted and/or terminated by one or more substituents selected from -OH, halo, cyano, nitro,  $-C(O)R^{10}$ ,  $-C(O)OR^{11}$ ,  $-N(H)S(O)_2R^{11a}$ ,  $C_{1-6}$  alkyl and/or  $C_{1-6}$  alkoxy) or, together with  $R^9$ , represents  $C_{3-7}$  alkylene;

$R^9$  represents H,  $C_{1-4}$  alkyl or, together with  $R^8$ , represents  $C_{3-7}$  alkylene;

- 15 Het<sup>1</sup> represents a five to twelve-membered heterocyclic ring containing one or more heteroatoms selected from oxygen, nitrogen and/or sulfur, and which also optionally includes one or more =O substituents;

$R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$  or  $R^{46}$  independently represent H or  $C_{1-3}$  alkyl;

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$R^5$  represents H, halo,  $C_{1-3}$  alkyl,  $-OR^{12}$ ,  $-N(R^{13})R^{12}$  or, together with  $R^6$ , represents =O;

$R^6$  represents H,  $C_{1-4}$  alkyl or, together with  $R^5$ , represents =O;

$R^{12}$  represents H,  $C_{1-6}$  alkyl,  $-S(O)_2-C_{1-4}$ -alkyl,  $-C(O)R^{14}$ ,  $-C(O)OR^{14}$ ,

- 25  $-C(O)N(R^{15})R^{15a}$  or aryl (which latter group is optionally substituted and/or terminated by one or more substituents selected from -OH, halo, cyano, nitro,  $-C(O)R^{10}$ ,  $-C(O)OR^{11}$ ,  $-N(H)S(O)_2R^{11a}$ ,  $C_{1-6}$  alkyl and/or  $C_{1-6}$  alkoxy);

$R^{13}$  represents H or  $C_{1-4}$  alkyl;

$R^{14}$  represents H or  $C_{1-6}$  alkyl;

$R^{15}$  and  $R^{15a}$  independently represent H or  $C_{1-4}$  alkyl, or together represent  $C_{3-6}$  alkylene, optionally interrupted by an O atom;

5 A represents a single bond,  $C_{1-6}$  alkylene,  $-N(R^{16})(CH_2)_r-$  or  $-O(CH_2)_r-$  (in which two latter groups, the  $-(CH_2)_r-$  group is attached to the bispidine nitrogen atom);

B represents a single bond,  $C_{1-4}$  alkylene,  $-(CH_2)_nN(R^{17})-$ ,  $-(CH_2)_nS(O)_p-$ ,  $-(CH_2)_nO-$  (in which three latter groups, the  $-(CH_2)_n-$  group is attached to  
10 the carbon atom bearing  $R^5$  and  $R^6$ ),  $-C(O)N(R^{17})-$  (in which latter group, the  $-C(O)-$  group is attached to the carbon atom bearing  $R^5$  and  $R^6$ ),  $-N(R^{17})C(O)O(CH_2)_n-$ ,  $-N(R^{17})(CH_2)_n-$  (in which two latter groups, the  $N(R^{17})$  group is attached to the carbon atom bearing  $R^5$  and  $R^6$ ) or  $-(CH_2)_mC(H)(OH)(CH_2)_n-$  (in which latter group, the  $-(CH_2)_m-$  group is  
15 attached to the carbon atom bearing  $R^5$  and  $R^6$ );

m represents 1, 2 or 3;

n and r independently represent 0, 1, 2, 3 or 4;

p represents 0, 1 or 2;

$R^{16}$  and  $R^{17}$  independently represent H or  $C_{1-4}$  alkyl;

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$R^7$  represents  $C_{1-6}$  alkyl, aryl or Het<sup>2</sup>, all of which groups are optionally substituted and/or terminated (as appropriate) by one or more substituents selected from -OH, cyano, halo, amino, nitro, Het<sup>3</sup>,  $-C(O)R^{10}$ ,  $-C(O)OR^{11}$ ,  $C_{1-6}$  alkyl,  $C_{1-6}$  alkoxy,  $-N(H)S(O)_2R^{18}$ ,  $-S(O)_2R^{19}$ ,  $-OS(O)_2R^{20}$ ,  
25  $-N(H)C(O)N(H)R^{21}$ ,  $-C(O)N(H)R^{22}$  and/or aryl (which latter group is optionally substituted by one or more cyano groups);

Het<sup>2</sup> and Het<sup>3</sup> independently represent a five to twelve-membered heterocyclic group containing one or more heteroatoms selected from

oxygen, nitrogen and/or sulfur, and which also optionally includes one or more =O substituents;

$R^{18}$ ,  $R^{19}$  and  $R^{20}$  independently represent  $C_{1-6}$  alkyl;

$R^{21}$  and  $R^{22}$  independently represent H or  $C_{1-6}$  alkyl (optionally terminated by cyano); and

$R^{10}$  and  $R^{11}$  independently represent, at each individual occurrence, H or  $C_{1-6}$  alkyl;

$R^{11a}$  represents, at each individual occurrence,  $C_{1-6}$  alkyl;

or a pharmaceutically acceptable derivative thereof;

provided that:

- (a) when A and B are both single bonds and  $R^7$  is optionally substituted aryl, then  $R^5$  and  $R^6$  do not both represent H;
- (b) when A represents a single bond, then  $R^5$  and  $R^6$  do not together represent =O; and
- (c) when  $R^5$  represents  $-OR^{12}$  or  $-N(R^{13})R^{12}$ , then:-
  - (i) A does not represent  $-N(R^{16})(CH_2)_r-$  or  $-O(CH_2)_r-$ ; and/or
  - (ii) n does not represent 0 when B represents  $-(CH_2)_nN(R^{17})-$ ,  $-(CH_2)_nS(O)_p-$  or  $-(CH_2)_nO-$ ,

which compounds are referred to hereinafter as "the compounds of the invention".

Aryl groups that may be mentioned include  $C_{6-10}$  aryl groups, such as phenyl, naphthyl and the like. Oxyaryl groups that may be mentioned include  $C_{6-10}$  oxyaryl groups, such as oxyphenyl (phenoxy), oxynaphthyl

(naphthoxy) and the like. When substituted, aryl and aryloxy groups are preferably substituted by one to three substituents.

Het<sup>1</sup>, Het<sup>2</sup> and Het<sup>3</sup> groups that may be mentioned include those  
 5 containing 1 to 4 heteroatoms (selected from the group oxygen, nitrogen  
 and/or sulfur) and in which the total number of atoms in the ring system  
 are between five and twelve. Het (Het<sup>1</sup>, Het<sup>2</sup> and Het<sup>3</sup>) groups may be  
 wholly/partly aromatic in character and may be bicyclic. Heterocyclic  
 groups that may be mentioned include morpholinyl, thiazolyl, oxazolyl,  
 10 isoxazolyl, cinnolinyl, quinazolinyl, phthalazinyl, purinyl, benzimidazolyl,  
 pyrimidinyl, piperazinyl, pyrazinyl, piperidinyl, pyridinyl, triazolyl,  
 imidazolyl, quinolinyl, isoquinolinyl, dioxanyl, benzodioxanyl,  
 benzodioxolyl, benzodioxepanyl, benzomorpholinyl, indolyl, pyrazolyl,  
 pyrrolyl, benzothiophenyl, thiophenyl, chromanyl, thiochromanyl,  
 15 benzofuranyl, pyranyl, tetrahydropyranyl, tetrahydrofuranyl, furanyl and  
 the like. Values of Het<sup>1</sup> that may be mentioned include tetrahydropyranyl,  
 isoxazolyl, benzodioxolyl, benzodioxepanyl and thiophenyl. Values of  
 Het<sup>2</sup> that may be mentioned include quinolinyl, isoquinolinyl,  
 benzomorpholinyl, benzodioxanyl, piperazinyl, indolyl and pyrazolyl.  
 20 Values of Het<sup>3</sup> that may be mentioned include imidazolyl. Substituents on  
 Het (Het<sup>1</sup>, Het<sup>2</sup> and Het<sup>3</sup>) groups may, where appropriate, be located on  
 any atom in the ring system including a heteroatom. The point of  
 attachment of Het (Het<sup>1</sup>, Het<sup>2</sup> and Het<sup>3</sup>) groups may be *via* any atom in the  
 ring system including (where appropriate) a heteroatom. Het (Het<sup>1</sup>, Het<sup>2</sup>  
 25 and Het<sup>3</sup>) groups may also be in the N- or S-oxidised form.

Pharmaceutically acceptable derivatives include salts and solvates. Salts  
 which may be mentioned include acid addition salts. Pharmaceutically  
 acceptable derivatives also include, at the bispidine nitrogens, C<sub>1-4</sub> alkyl

quaternary ammonium salts and N-oxides, provided that when a N-oxide is present:

- (a) no Het (Het<sup>1</sup>, Het<sup>2</sup>, Het<sup>3</sup>) group contains an unoxidised S-atom; and/or
- (b) p does not represent 0 when B represents  $-(CH_2)_nS(O)_p^-$ .

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The compounds of the invention may exhibit tautomerism. All tautomeric forms and mixtures thereof are included within the scope of the invention.

The compounds of the invention may also contain one or more asymmetric carbon atoms and may therefore exhibit optical and/or diastereoisomerism. Diastereoisomers may be separated using conventional techniques, e.g. chromatography or fractional crystallisation. The various stereoisomers may be isolated by separation of a racemic or other mixture of the compounds using conventional, e.g. fractional crystallisation or HPLC, techniques.

Alternatively the desired optical isomers may be made by reaction of the appropriate optically active starting materials under conditions which will not cause racemisation or epimerisation, or by derivatisation, for example with a homochiral acid followed by separation of the diastereomeric esters by conventional means (e.g. HPLC, chromatography over silica). All stereoisomers are included within the scope of the invention.

Alkyl groups that R<sup>1</sup>, R<sup>2</sup>, R<sup>2b</sup>, R<sup>2c</sup>, R<sup>2d</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup>, R<sup>11</sup>, R<sup>11a</sup>, R<sup>12</sup>, R<sup>13</sup>, R<sup>14</sup>, R<sup>15</sup>, R<sup>15a</sup>, R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup>, R<sup>19</sup>, R<sup>20</sup>, R<sup>21</sup>, R<sup>22</sup>, R<sup>41</sup>, R<sup>42</sup>, R<sup>43</sup>, R<sup>44</sup>, R<sup>45</sup> and R<sup>46</sup> may represent, that R<sup>12</sup> may include, and with which R<sup>3</sup>, R<sup>4</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>12</sup> may be substituted; and alkoxy groups that R<sup>4</sup> may represent, and with which R<sup>4</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>12</sup> may be substituted; may be linear or, when there is a sufficient number (i.e. three) of carbon atoms, be branched and/or cyclic. Further, when there is a sufficient number (i.e. four) of carbon atoms, such alkyl and alkoxy groups may



also be part cyclic/acyclic. Such alkyl and alkoxy groups may also be saturated or, when there is a sufficient number (i.e. two) of carbon atoms, be unsaturated and/or interrupted by oxygen.

- 5 Alkylene groups that  $R^3$  and  $R^4$ ,  $R^8$  and  $R^9$ ,  $R^{15}$  and  $R^{15a}$ , A, and B, may represent; and  $-(CH_2)_m-$ ,  $-(CH_2)_n-$ ,  $-(CH_2)_q-$  and  $-(CH_2)_r-$  chains that A, B and  $R^4$  (as appropriate) may include, may be linear or, when there is a sufficient number (i.e. two) of carbon atoms, be branched. Such alkylene groups and  $-(CH_2)-$  containing chains may also be saturated or, when there  
10 is a sufficient number (i.e. two) of carbon atoms, be unsaturated and/or interrupted by oxygen.

Halo groups that  $R^5$  may represent, and with which  $R^4$ ,  $R^7$ ,  $R^8$  and  $R^{12}$  may be substituted, include fluoro, chloro, bromo and iodo.

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- For the avoidance of doubt, each  $R^{10}$ ,  $R^{11}$ , and  $R^{11a}$ , group identified herein is independent of other  $R^{10}$ ,  $R^{11}$ , and  $R^{11a}$ , groups, respectively. For example, when  $R^4$  and  $R^7$  both represent aryl substituted by  $-C(O)R^{10}$ , the two individual  $-C(O)R^{10}$  substituents are independent of one another, and are  
20 not necessarily identical (though this possibility is not excluded).

Abbreviations are listed at the end of this specification.

- According to a further aspect of the invention there is provided  
25 compounds of formula I as hereinbefore defined, but with the further provisos that:

- (a) when A represents  $-N(R^{16})(CH_2)_r-$  or  $-O(CH_2)_r-$ , then r does not represent 0 or 1; and

- (b) when  $R^5$  represents  $-OH$  or  $-N(R^{13})R^{12}$ , then B does not represent  $-N(R^{17})C(O)O(CH_2)_n-$  or  $-N(R^{17})(CH_2)_n-$ .

Preferred compounds of the invention include those in which:

5  $R^1$  represents H;

$R^2$  represents H;

$R^3$  represents

H;

$C_{1-2}$  alkyl; or,

10 together with  $R^4$  represents  $C_{4-5}$  alkylene, optionally interrupted by an O atom and/or optionally substituted by one or more methyl groups;

$R^4$  represents

H;

15 linear or branched and/or saturated or unsaturated and/or cyclic, acyclic and/or part cyclic/acyclic  $C_{1-8}$  alkyl (which alkyl group is optionally substituted by one or more cyano or halo groups and/or interrupted by an O atom);

$C_{1-6}$  alkoxy;

$-(CH_2)_qS(O)_2R^8$ ,  $-(CH_2)_qC(O)OR^8$ ,  $-(CH_2)_qN(H)C(O)R^8$ ,

20  $-(CH_2)_qC(O)R^8$ , (in which latter four groups, q represents 0, 1 or 2 and  $R^8$  represents linear or branched and/or acyclic, cyclic and/or part cyclic/acyclic  $C_{1-4}$  alkyl, or phenyl (which phenyl group is optionally substituted by one or more cyano and/or  $C_{1-3}$  alkyl groups));

25  $-(CH_2)_qC(O)N(R^9)R^8$  (in which latter group, q represents 0, 1 or 2 and  $R^8$  and  $R^9$  independently represent H, linear or branched and/or acyclic, cyclic and/or part cyclic/acyclic  $C_{1-4}$  alkyl, or together represent  $C_{4-6}$  alkylene);

$-(CH_2)_q$ -phenyl,  $-(CH_2)_q$ -oxyphenyl or  $-(CH_2)_q$ -Het<sup>1</sup> (in which latter three groups, q represents 0, 1, 2 or 3, the  $-(CH_2)_q$ - part is optionally

substituted by a cyano group, and the phenyl, or Het<sup>1</sup>, part is optionally substituted with one or more substituents selected from cyano, nitro, linear or branched C<sub>1-4</sub> alkyl, linear or branched C<sub>1-4</sub> alkoxy and N(H)S(O)<sub>2</sub>R<sup>11a</sup>); or,

5 together with R<sup>3</sup>, represents C<sub>4-5</sub> alkylene, optionally interrupted by an O atom and/or optionally substituted by one or more methyl groups;

R<sup>5</sup> represents

H;

fluoro;

10 OR<sup>12</sup> (in which R<sup>12</sup> represents H, phenyl (optionally substituted by one or more methoxy groups) or C(O)N(H)R<sup>15a</sup> (in which R<sup>15a</sup> represents linear or branched C<sub>1-4</sub> alkyl));

-N(R<sup>13</sup>)(R<sup>12</sup>) (in which R<sup>12</sup> represents H, C<sub>1-2</sub> alkyl, -S(O)<sub>2</sub>-C<sub>1-2</sub> alkyl, -C(O)R<sup>14</sup> (in which R<sup>14</sup> represents C<sub>1-2</sub> alkyl), -C(O)OR<sup>14</sup> (in which R<sup>14</sup> represents linear or branched C<sub>1-5</sub> alkyl) or -C(O)N(R<sup>15</sup>)(R<sup>15a</sup>) (in which R<sup>15</sup> and R<sup>15a</sup> independently represent H or linear or branched C<sub>1-3</sub> alkyl or together represent C<sub>4-5</sub> alkylene, which alkylene group is optionally interrupted by an O atom) and R<sup>13</sup> represents H or C<sub>1-2</sub> alkyl); or,

20 together with R<sup>6</sup>, represents =O (especially in the case where R<sup>7</sup> represents alkyl or Het<sup>2</sup>);

R<sup>6</sup> represents H or C<sub>1-2</sub> alkyl or together with R<sup>5</sup> represents =O (especially in the case where R<sup>7</sup> represents alkyl or Het<sup>2</sup>);

A represents a single bond, linear or branched C<sub>1-4</sub> alkylene (which group is also optionally interrupted by O), -N(H)(CH<sub>2</sub>)<sub>r</sub>- or -O(CH<sub>2</sub>)<sub>r</sub>- (in which 25 latter two groups r is 1 or 2);

B represents a single bond, C<sub>1-4</sub> alkylene, -(CH<sub>2</sub>)<sub>n</sub>O-, -(CH<sub>2</sub>)<sub>n</sub>S(O)<sub>2</sub>-, -(CH<sub>2</sub>)<sub>n</sub>N(H)- or -N(H)(CH<sub>2</sub>)<sub>n</sub>- (in which latter four cases n is 0, 1, 2 or 3);

R<sup>7</sup> represents

linear or branched and/or acyclic, cyclic and/or part cyclic/acyclic  $C_{1-6}$  alkyl (optionally substituted and/or terminated by OH);

Het<sup>2</sup> (optionally substituted by one or more substituents selected from cyano,  $C_{1-3}$  alkyl, phenyl (which latter group is optionally substituted with one or more cyano groups), =O,  $C(O)R^{10}$  (in which  $R^{10}$  is linear or branched  $C_{1-3}$  alkyl) or  $S(O)_2R^{19}$  (in which  $R^{19}$  is  $C_{1-2}$  alkyl)); or

phenyl (optionally substituted by one or more substituents selected from cyano, nitro, linear or branched  $C_{1-3}$  alkyl, linear or branched  $C_{1-3}$  alkoxy, fluoro, chloro,  $C(O)N(H)R^{22}$  (in which  $R^{22}$  represents linear or branched and/or acyclic, cyclic and/or part cyclic/acyclic  $C_{1-4}$  alkyl, which alkyl group is optionally terminated by cyano),  $N(H)S(O)_2R^{18}$  (in which  $R^{18}$  represents  $C_{1-2}$  alkyl) or Het<sup>3</sup>);  
 $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$  and  $R^{46}$  all represent H.

More preferred compounds of the invention include those in which:

$R^3$  represents H;

$R^5$  represents H, OH or  $-N(H)C(O)N(R^{15})(R^{15a})$ ;

$R^6$  represents H;

A represents  $-CH_2-$  or  $-(CH_2)_2-$ ;

B represents a single bond,  $-CH_2N(H)-$  or  $-CH_2O-$  (where, for the avoidance of doubt, the  $-CH_2-$  part is attached to the carbon atom bearing  $R^5$  and  $R^6$ );

$R^7$  represents phenyl (substituted by a cyano group (preferably in the 4-position relative to B) and by one or more optional  $C(O)N(H)R^{22}$  substituent).

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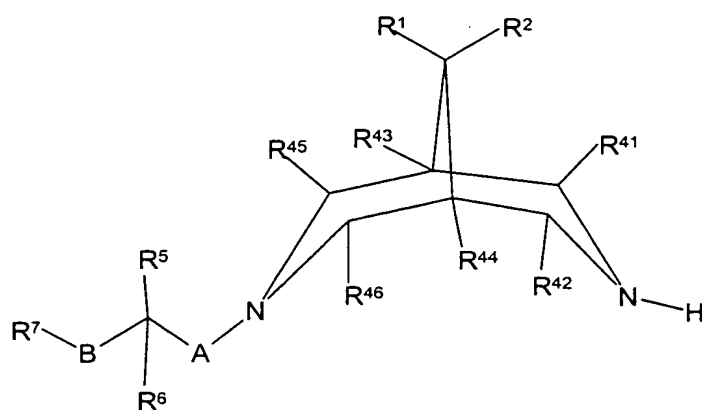
Preferred compounds of the invention include the compounds of the Examples disclosed hereinafter.

## Preparation

According to the invention there is also provided a process for the preparation of compounds of formula I which comprises:

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(a) for compounds of formula I in which  $R^3$  is H, reaction of a compound of formula II,



II

10

wherein  $R^1$ ,  $R^2$ ,  $R^5$ ,  $R^6$ ,  $R^7$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$ ,  $R^{46}$ , A and B are as hereinbefore defined with a compound of formula III,



15 wherein  $R^4$  is as hereinbefore defined, for example at between  $0^\circ\text{C}$  and reflux temperature in the presence of an appropriate organic solvent (e.g. dichloromethane), or *via* solid phase synthesis under conditions known to those skilled in the art;

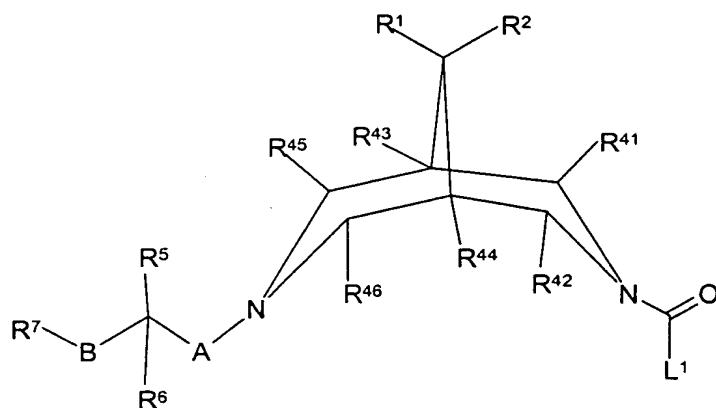
20 (b) reaction of a compound of formula II, as hereinbefore defined, with a carbonic acid derivative of formula IV,



wherein  $L^1$  represents a leaving group such as halo, imidazole or  $R^{23}O$ -  
 (wherein  $R^{23}$  represents, for example,  $C_{1-10}$  alkyl, aryl or  $C_{1-3}$  alkylaryl,  
 which groups are optionally substituted by one or more halo or nitro groups)  
 and  $R^3$  and  $R^4$  are as hereinbefore defined, for example at between room  
 5 and reflux temperature in the presence of a suitable base (e.g. triethylamine  
 or potassium carbonate) and an appropriate organic solvent (e.g.  
 dichloromethane, THF, acetonitrile, toluene, or mixtures thereof);

(c) reaction of a compound of formula V,

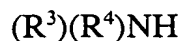
10



V

wherein  $R^1$ ,  $R^2$ ,  $R^5$ ,  $R^6$ ,  $R^7$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$ ,  $R^{46}$ , A, B and  $L^1$  are as  
 hereinbefore defined with a compound of formula VA,

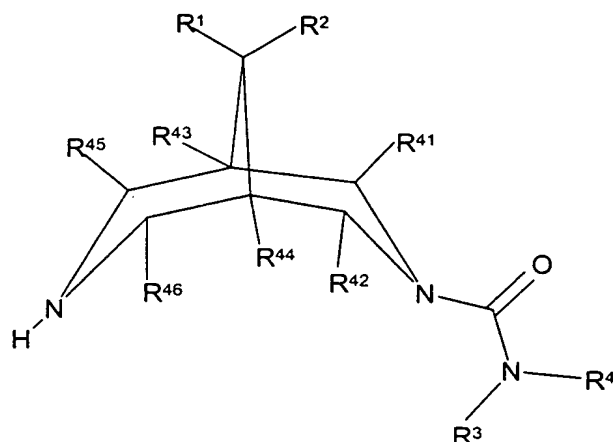
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VA

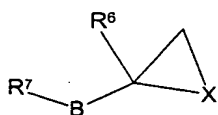
wherein  $R^3$  and  $R^4$  are as hereinbefore defined, for example at between  
 room and reflux temperature in the presence of a suitable base (e.g.  
 triethylamine or potassium carbonate) and an appropriate organic solvent  
 (e.g. dichloromethane, THF, acetonitrile, toluene, or mixtures thereof), or  
 20 *via* solid phase synthesis under conditions known to those skilled in the art;

(d) for compounds of formula I in which A represents  $\text{CH}_2$  and  $\text{R}^5$  represents  $-\text{OH}$  or  $-\text{N}(\text{H})\text{R}^{12}$ , wherein  $\text{R}^{12}$  is as hereinbefore defined, reaction of a compound of formula VI,



VI

wherein  $\text{R}^1$ ,  $\text{R}^2$ ,  $\text{R}^3$ ,  $\text{R}^4$ ,  $\text{R}^{41}$ ,  $\text{R}^{42}$ ,  $\text{R}^{43}$ ,  $\text{R}^{44}$ ,  $\text{R}^{45}$  and  $\text{R}^{46}$  are as hereinbefore defined, with a compound of formula VII,



VII

wherein X represents O or  $\text{N}(\text{R}^{12})$  and  $\text{R}^6$ ,  $\text{R}^7$ ,  $\text{R}^{12}$  and B are as hereinbefore defined, for example at elevated temperature (e.g.  $60^\circ\text{C}$  to reflux) in the presence of a suitable solvent (e.g. a lower alkyl alcohol (e.g. IPA), acetonitrile, or a mixture of a lower alkyl alcohol and water);

(e) reaction of a compound of formula VI, as hereinbefore defined, with a compound of formula VIII,

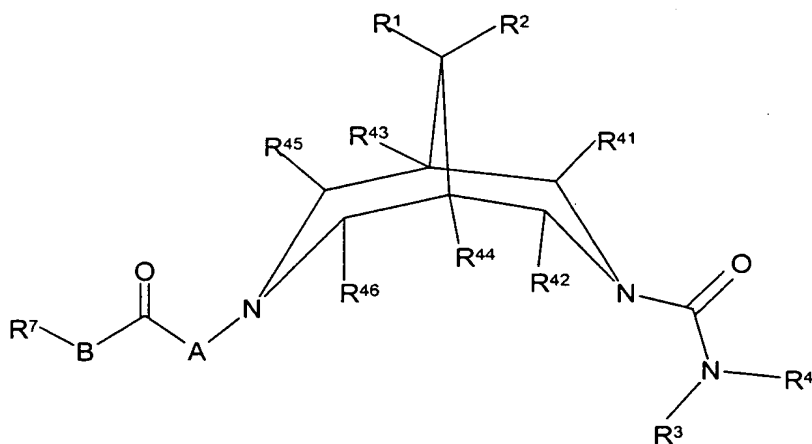
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VIII

wherein  $L^2$  represents a leaving group (e.g. mesylate, tosylate or halo) and  $R^5$ ,  $R^6$ ,  $R^7$ , A and B are as hereinbefore defined, for example at elevated temperature (e.g. between  $35^\circ\text{C}$  and reflux temperature) in the presence of a suitable base (e.g. triethylamine or  $\text{K}_2\text{CO}_3$ ) and an appropriate organic solvent (e.g. acetonitrile or DMSO);

(f) for compounds of formula I in which  $R^5$  represents H or OH and  $R^6$  represents H, reduction of a compound of formula IX,



IX

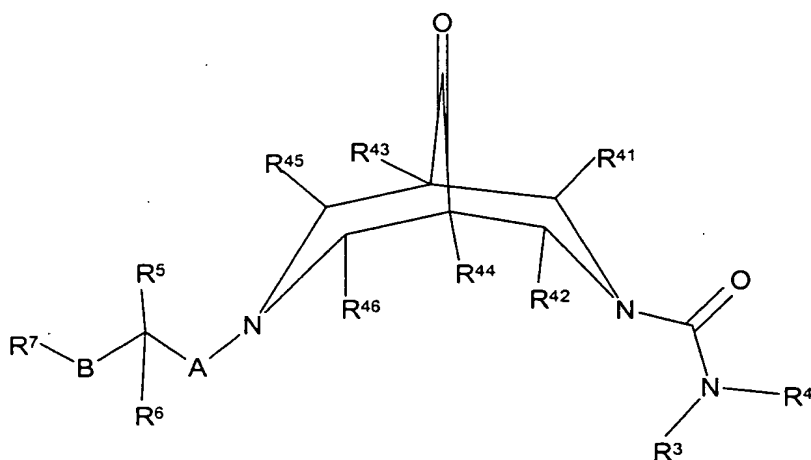
wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ ,  $R^7$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$ ,  $R^{46}$ , A and B are as hereinbefore defined, in the presence of a suitable reducing agent and under appropriate reaction conditions; for example, for formation of compounds of formula I in which  $R^5$  represents OH, reduction may be performed under mild reaction conditions in the presence of e.g. sodium borohydride and an appropriate organic solvent (e.g. THF); and for formation of compounds of formula I in which  $R^5$  represents H, reduction may be performed by



activating the relevant C=O group using an appropriate agent (such as tosylhydrazine) in the presence of a suitable reducing agent (e.g. sodium borohydride or sodium cyanoborohydride) and an appropriate organic solvent (e.g. a lower (e.g. C<sub>1-6</sub>) alkyl alcohol);

5

(g) for compounds of formula I in which R<sup>1</sup> and R<sup>2</sup> both represent H, reduction of a corresponding compound of formula X,



X

10

wherein R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>41</sup>, R<sup>42</sup>, R<sup>43</sup>, R<sup>44</sup>, R<sup>45</sup>, R<sup>46</sup>, A and B are as hereinbefore defined, and in which the bridgehead C=O group may be activated using an appropriate agent, such as tosylhydrazine, in the presence of a suitable reducing agent (e.g. sodium borohydride, sodium cyanoborohydride) and an appropriate organic solvent (e.g. a lower alkyl alcohol), or under standard Wolff-Kischner conditions known to those skilled in the art; when the C=O group is activated, the activation step may be carried out at between room and reflux temperature in the presence of an appropriate organic solvent (e.g. a lower alkyl alcohol such as methanol, ethanol or IPA), whereafter the reducing agent may be added to the reaction

20

(h) for compounds of formula I in which R<sup>1</sup> and R<sup>2</sup> together represent -O(CH<sub>2</sub>)<sub>2</sub>O-, reaction of a corresponding compound of formula X as hereinbefore defined with ethane-1,2-diol under appropriate reaction conditions, for example by refluxing in the presence of *p*TSA and an appropriate organic solvent (e.g. toluene);

The chemical structure shows a bicyclic system, specifically a decalin derivative, with various substituents. On the left, a hydroxyalkyl group is attached to one of the bridgehead carbons:  $\text{HO}-(\text{CH}_2)_n-\text{C}(\text{R}^5)(\text{R}^6)-\text{A}-\text{N}$ , where the nitrogen is part of the bicyclic ring. The bridgehead carbons are also substituted with  $\text{R}^{45}$  and  $\text{R}^{46}$ . The other bridgehead carbon is part of a carbamate group:  $-\text{N}-\text{C}(=\text{O})-\text{N}(\text{R}^3)(\text{R}^4)$ . The bicyclic system has several other substituents:  $\text{R}^{41}$  and  $\text{R}^{42}$  on the ring carbons,  $\text{R}^{43}$  and  $\text{R}^{44}$  on the bridge carbons, and  $\text{R}^1$  and  $\text{R}^2$  on the carbons at the top of the bicyclic system.

wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>41</sup>, R<sup>42</sup>, R<sup>43</sup>, R<sup>44</sup>, R<sup>45</sup>, R<sup>46</sup>, A and n are as hereinbefore defined, with a compound of formula XIA,

XIA

in which R<sup>7</sup> is as hereinbefore defined, for example under Mitsunobu-type conditions e.g. at between ambient (e.g. 25°C) and reflux temperature in the presence of a tertiary phosphine (e.g. tributylphosphine or triphenylphosphine), an azodicarboxylate derivative (e.g. diethylazodicarboxylate or 1,1'-(azodicarbonyl)dipiperidine) and an appropriate organic solvent (e.g. dichloromethane or toluene);

(j) for compounds of formula I which are bispidine-nitrogen N-oxide derivatives, oxidation of the corresponding bispidine nitrogen of a corresponding compound of formula I, in the presence of a suitable oxidising agent (e.g. *m*CPBA), for example at 0°C in the presence of a suitable organic solvent (e.g. DCM);

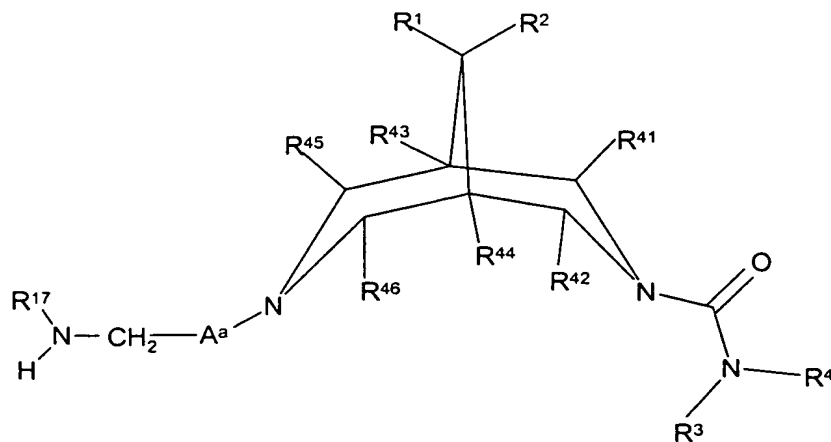
(k) for compounds of formula I which are C<sub>1-4</sub> alkyl quaternary ammonium salt derivatives, in which the alkyl group is attached to a bispidine nitrogen, reaction, at the bispidine nitrogen, of a corresponding compound of formula I with a compound of formula XII,



XII

wherein R<sup>b</sup> represents C<sub>1-4</sub> alkyl and L<sup>3</sup> is a leaving group such as halo, alkane sulfonate or aryl sulfonate, for example at room temperature in the presence of an appropriate organic solvent (e.g. DMF), followed by purification (using e.g. HPLC) in the presence of a suitable counter-ion provider (e.g. NH<sub>4</sub>OAc);

(l) for compounds of formula I in which R<sup>5</sup> and R<sup>6</sup> represent H, A represents C<sub>1-6</sub> alkylene and B represents -N(R<sup>17</sup>)(CH<sub>2</sub>)<sub>n</sub>-, reaction of a compound of formula XIII,



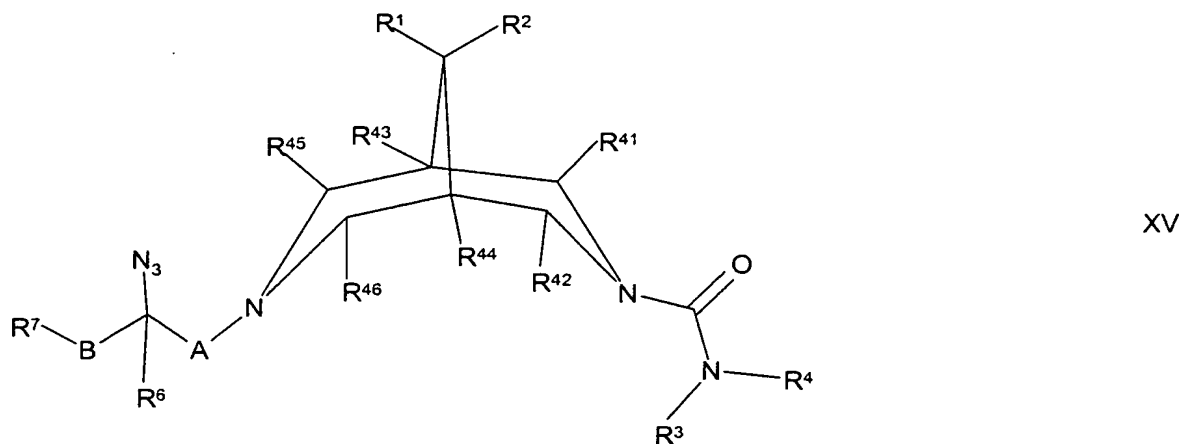
XIII

wherein A<sup>a</sup> represents C<sub>1-6</sub> alkylene and R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>41</sup>, R<sup>42</sup>, R<sup>43</sup>, R<sup>44</sup>, R<sup>45</sup>, R<sup>46</sup> and R<sup>17</sup> are as hereinbefore defined with a compound of formula XIV,



wherein R<sup>7</sup>, n and L<sup>2</sup> are as hereinbefore defined, for example at 40°C in the presence of a suitable organic solvent (e.g. acetonitrile);

(m) for compounds of formula I in which R<sup>5</sup> represents -NH<sub>2</sub>, reduction of a corresponding compound of formula XV,



wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>41</sup>, R<sup>42</sup>, R<sup>43</sup>, R<sup>44</sup>, R<sup>45</sup>, R<sup>46</sup>, A and B are as hereinbefore defined, for example by hydrogenation at a suitable pressure in the presence of a suitable catalyst (e.g. palladium on carbon) and an appropriate solvent (e.g. a water-ethanol mixture);

(n) for compounds of formula I in which R<sup>5</sup> represents -N(R<sup>13</sup>)C(O)NH(R<sup>15</sup>), reaction of a corresponding compound of formula I in which R<sup>5</sup> represents -N(R<sup>13</sup>)H with a compound of formula XVI,



wherein R<sup>15</sup> is as hereinbefore defined, for example at ambient temperature (e.g. 25°C) in the presence of a suitable solvent (e.g. benzene);

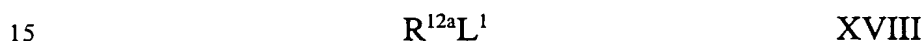
(o) for compounds of formula I in which  $R^5$  represents  $-N(R^{13})C(O)R^{14}$ , reaction of a corresponding compound of formula I in which  $R^5$  represents  $-N(R^{13})H$  with a compound of formula XVII,



5 wherein  $R^x$  represents a suitable leaving group, such as  $C_{1-4}$  alkoxy, halo (e.g. Cl, Br) or *p*-nitrophenyl, and  $R^{14}$  is as hereinbefore defined, for example at between ambient and reflux temperature in the presence of a suitable solvent (e.g. dichloromethane or acetonitrile) and optionally in the presence of a suitable base (e.g. triethylamine or potassium carbonate);

10

(p) for compounds of formula I in which  $R^5$  represents  $-N(H)R^{12}$ , wherein  $R^{12}$  is as previously defined provided that it does not represent H, reaction of a corresponding compound of formula I, in which  $R^5$  represents  $-NH_2$  with a compound of formula XVIII,



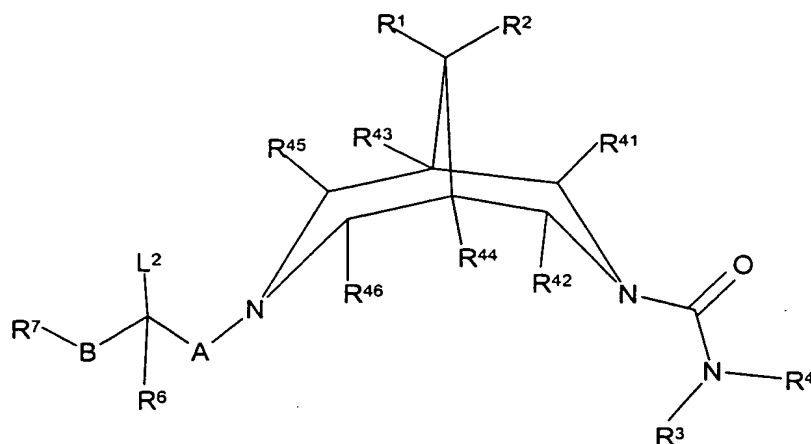
15 wherein  $R^{12a}$  represents  $R^{12}$  as hereinbefore defined except that it does not represent H and  $L^1$  is as hereinbefore defined, for example under conditions that are well known to those skilled in the art;

20 (q) for compounds of formula I in which  $R^5$  represents  $-OR^{12}$  in which  $R^{12}$  represents  $C_{1-6}$  alkyl or optionally substituted aryl, reaction of a corresponding compound of formula I in which  $R^5$  represents  $-OH$  with a compound of formula XIX,



25 wherein  $R^{12a}$  represents  $C_{1-6}$  alkyl or optionally substituted aryl, for example at between ambient (e.g. 25°C) and reflux temperature, under Mitsunobu-type conditions (i.e. in the presence of e.g. triphenylphosphine, an azodicarboxylate derivative (e.g. 1,1'-(azodicarbonyl)dipiperidine) and a suitable organic solvent (e.g. dichloromethane));

(r) for compounds of formula I in which  $R^5$  represents  $-OR^{12}$ , in which  $R^{12}$  represents  $C_{1-6}$  alkyl or optionally substituted aryl, reaction of a compound of formula XX,



XX

wherein  $L^2$ ,  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ ,  $R^6$ ,  $R^7$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$ ,  $R^{46}$ , A and B are as hereinbefore defined with a compound of formula XIX as hereinbefore defined, for example at between ambient (e.g.  $25^\circ\text{C}$ ) and reflux temperature, under Williamson-type conditions (i.e. in the presence of an appropriate base (e.g. KOH or NaH) and a suitable organic solvent (e.g. dimethylsulfoxide or DMF));

(s) for compounds of formula I in which  $R^5$  represents  $OR^{12}$  and  $R^{12}$  represents  $C(O)R^{14}$  and  $R^{14}$  is as hereinbefore defined, reaction of a corresponding compound of formula I as hereinbefore defined in which  $R^5$  represents OH with a compound of formula XXI,



XXI

wherein  $R^{14}$  is as hereinbefore defined, for example at ambient temperature (e.g.  $25^\circ\text{C}$ ) in the presence of a suitable coupling agent (e.g. 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide), an appropriate catalyst (e.g. 4-dimethylaminopyridine) and a reaction-inert organic solvent (e.g. THF);

(t) for compounds of formula I in which  $R^5$  represents halo, substitution of a corresponding compound of formula I in which  $R^5$  represents  $-OH$ , using an appropriate halogenating agent (e.g., for compounds in which  $R^5$  represents fluoro, reaction with diethylaminosulfurtrifluoride);

5

(u) for compounds of formula I in which  $R^3$  and/or  $R^4$  as appropriate represent alkyl groups (e.g.  $C_{1-6}$  or  $C_{1-12}$  alkyl, as appropriate), alkylation of a corresponding compound of formula I, in which  $R^3$  and/or  $R^4$  (as appropriate) represent H under conditions well known to those skilled in the art;

10

(v) conversion of one  $R^4$  group to another (e.g. conversion of  $-(CH_2)_qC(O)OR^8$  to  $-(CH_2)_qC(O)N(R^9)R^8$ , wherein  $R^8$ ,  $R^9$  and  $q$  are as hereinbefore defined) using techniques well known to those skilled in the art; or

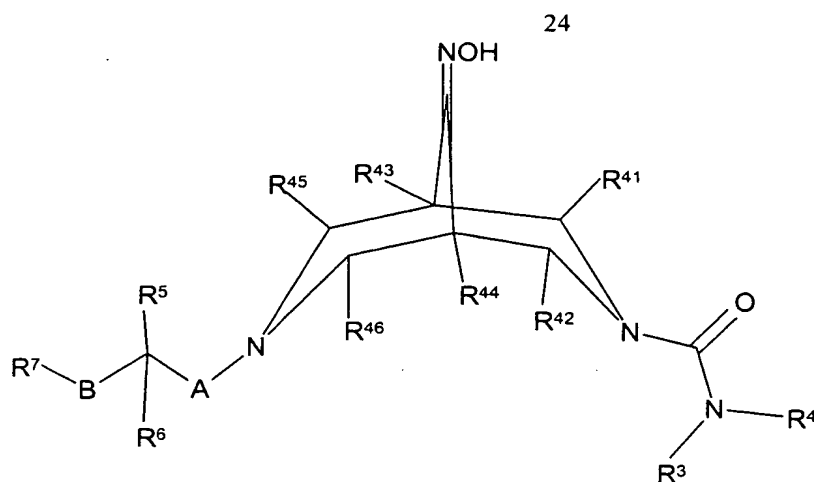
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(w) for compounds of formula I in which one of  $R^1$  and  $R^2$  represents H, and the other represents  $-OH$ , reduction of a corresponding compound of formula X, as hereinbefore defined, in the presence of a mild reducing agent, e.g. sodium borohydride, and an appropriate organic solvent (e.g. a lower alcohol such as methanol or ethanol);

20

(x) for compounds of formula I in which one of  $R^2$  and  $R^3$  represents  $-NH_2$  and the other represents H, reduction of a compound of formula XXIA,

25



wherein  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$ ,  $R^7$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$ ,  $R^{46}$ , A and B are as  
 5 hereinbefore defined, in the presence of a suitable reducing agent (e.g.  $\text{LiAlH}_4$ ), for example under conditions that are well known to those skilled in the art;

(y) for compounds of formula I in which one or both of  $R^1$  and  $R^2$   
 10 represent  $-\text{N}(\text{R}^{2c})\text{R}^{2d}$  in which one or both of  $\text{R}^{2c}$  and  $\text{R}^{2d}$  represents  $\text{C}_{1-6}$  alkyl, alkylation of a corresponding compound of formula I in which  $\text{R}^1$  and/or  $\text{R}^2$  represent  $-\text{N}(\text{R}^{2c})\text{R}^{2d}$  (as appropriate) in which  $\text{R}^{2c}$  and/or  $\text{R}^{2d}$  (as appropriate) represent H, using a compound of formula XXIB,



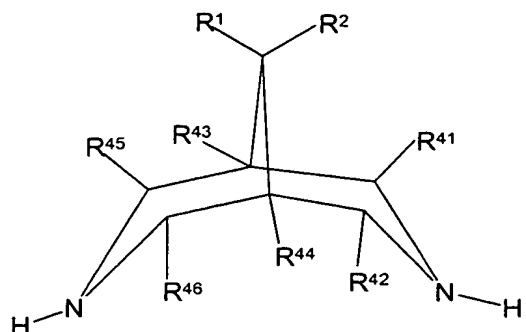
XXIB

15 wherein  $\text{R}^{2e}$  represents  $\text{C}_{1-6}$  alkyl and  $\text{L}^1$  is as hereinbefore defined, for example under conditions that are well known to those skilled in the art; or

(z) conversion of one substituent on  $\text{R}^7$  to another using techniques well  
 20 known to those skilled in the art.



Compounds of formula II may be prepared by reaction of a compound of formula XXII,



XXII

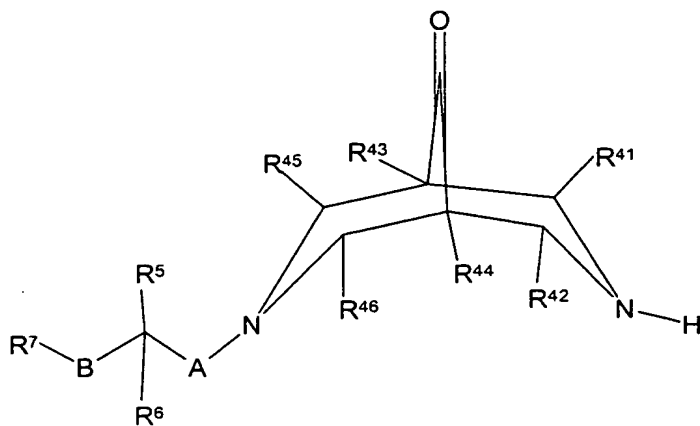
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wherein  $R^1$ ,  $R^2$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$  and  $R^{46}$  are as hereinbefore defined, with a compound of formula VIII as hereinbefore defined, for example as described hereinbefore for synthesis of compounds of formula I (process step (e)), or, in the case of compounds of formula II wherein A represents  $CH_2$  and  $R^5$  represents OH or  $N(H)R^{12}$ , with a compound of formula VII as hereinbefore defined, for example as described hereinbefore for synthesis of compounds of formula I (process step (d)).

10

15

Compounds of formula II in which  $R^1$  and  $R^2$  both represent H may be prepared by reduction of a compound of formula XXIII,



XXIII

wherein  $R^5$ ,  $R^6$ ,  $R^7$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$ ,  $R^{46}$ , A and B are as hereinbefore defined, and in which the C=O group may be activated using an appropriate agent, such as tosylhydrazine, for example as described hereinbefore for synthesis of compounds of formula I (process step (g)).

5

Compounds of formula IV may be prepared by reaction of a compound of formula VA, as hereinbefore defined, with a compound of formula XXIV,



wherein  $L^1$  is as hereinbefore defined, and in which the two  $L^1$  groups may be the same or different, for example at between 0°C and reflux temperature in the presence of a suitable base (e.g. triethylamine or potassium carbonate) and an appropriate organic solvent (e.g. toluene or dichloromethane).

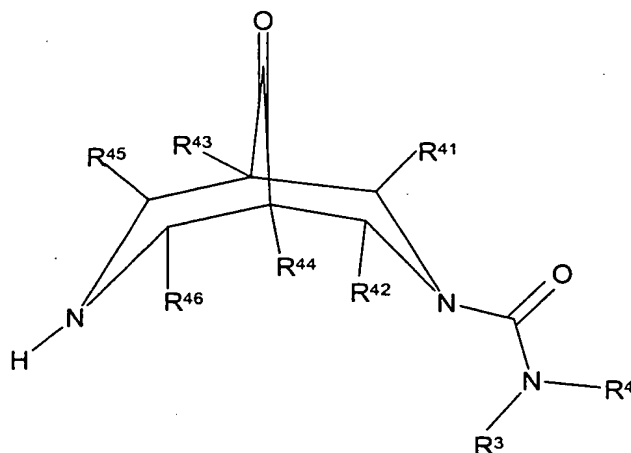
Compounds of formula V may be prepared by reaction of a compound of formula II, as hereinbefore defined, with a compound of formula XXIV, as hereinbefore defined, for example as described hereinbefore for the synthesis of compounds of formula IV.

Compounds of formula VI may be prepared by reaction of a compound of formula XXII, as hereinbefore defined, with a compound of formula III, as hereinbefore defined, for example as described hereinbefore for synthesis of compounds of formula I (process step (a)), or with a compound of formula IV, as hereinbefore defined, for example as described hereinbefore for synthesis of compounds of formula I (process step (b)).

Compounds of formula VI may alternatively be prepared by reaction of a compound of formula XXII, as hereinbefore defined, with a compound of formula XXIV, as hereinbefore defined, for example as described

hereinbefore for synthesis of compounds of formula IV, followed by reaction of the resultant intermediate with a compound of formula VA, as hereinbefore defined, for example as described hereinbefore for the synthesis of compounds of formula I (process step (c)).

Compounds of formula VI in which  $R^1$  and  $R^2$  represent H may alternatively be prepared by reduction of a corresponding compound of formula XXV,



XXV

wherein  $R^3$ ,  $R^4$ ,  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$  and  $R^{46}$  are as hereinbefore defined, and in which the C=O group may be activated using an appropriate agent, such as tosylhydrazine, for example as described hereinbefore for compounds of formula I (process step (g)).

Compounds of formula VI in which one or more of  $R^{41}$ ,  $R^{42}$ ,  $R^{45}$  and/or  $R^{46}$  represent  $C_{1-3}$  alkyl may be prepared by reaction of a compound of formula VI in which  $R^{41}$ ,  $R^{42}$ ,  $R^{45}$  and/or  $R^{46}$  (as appropriate) represent H, with an appropriate alkylating agent (e.g. dimethyl sulfate), for example in the presence of a suitable strong base (e.g. *s*-BuLi), N,N,N',N'-tetramethylethylenediamine and a reaction-inert solvent (e.g. THF).

Compounds of formula VII may be prepared in accordance with techniques which are known to those skilled in the art. For example, compounds of formula VII in which:

- 5 (1) B represents  $-\text{CH}_2\text{O}-$  and X represents O may be prepared by reaction of a compound of formula XIA as hereinbefore defined, with a compound of formula XXVI,

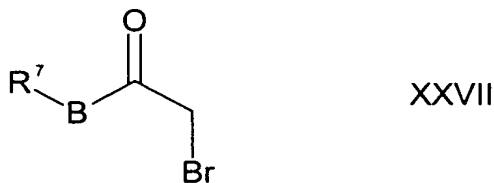


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wherein  $\text{R}^6$  and  $\text{L}^2$  are as hereinbefore defined, for example at elevated temperature (e.g. between  $60^\circ\text{C}$  and reflux temperature) in the presence of a suitable base (e.g.  $\text{K}_2\text{CO}_3$  or  $\text{NaOH}$ ) and an appropriate organic solvent (e.g. acetonitrile or toluene/water), or as otherwise described in the prior art;

15

- (2)  $\text{R}^6$  represents H and X represents O may be prepared by reduction of a compound of formula XXVII,



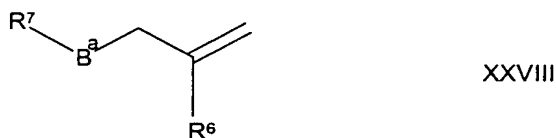
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wherein  $\text{R}^7$  and B are as hereinbefore defined, for example at between  $-15^\circ\text{C}$  and room temperature in the presence of a suitable reducing agent (e.g.  $\text{NaBH}_4$ ) and an appropriate organic solvent (e.g. THF), followed by

25 an internal displacement reaction in the resultant intermediate, for example

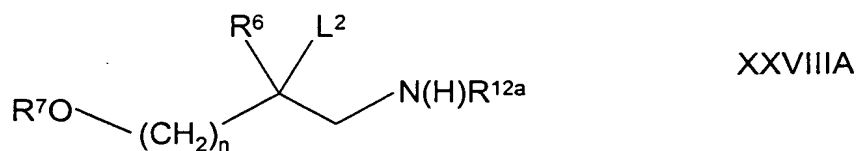
at room temperature in the presence of a suitable base (e.g.  $K_2CO_3$ ) and an appropriate organic solvent (e.g. acetonitrile);

- (3) B represents  $C_{1-4}$  alkylene,  $-(CH_2)_nN(R^{17})-$ ,  $-(CH_2)_nS(O)_2-$  or  $-(CH_2)_nO-$  (in which latter three groups n represents 1, 2, 3 or 4) or  $-(CH_2)_mC(H)(OH)(CH_2)_n-$  and X represents O may be prepared by oxidation of a compound of formula XXVIII,



- in which  $B^a$  represents a single bond,  $C_{1-3}$  alkylene,  $-(CH_2)_nN(R^{17})-$ ,  $-(CH_2)_nS(O)_2-$  or  $-(CH_2)_nO-$  (in which latter three groups n represents 1, 2, 3 or 4) or  $-(CH_2)_mC(H)(OH)(CH_2)_n-$  (in which latter group n is as hereinbefore defined), and in all cases  $R^{17}$  and m are as hereinbefore defined, in the presence of a suitable oxidising agent (e.g. *m*CPBA), for example by refluxing in the presence of a suitable organic solvent (e.g. DCM); or

- (4) B represents  $-(CH_2)_nO-$  and X represents  $N(R^{12})$  and  $R^{12}$  represents  $-S(O)_2-C_{1-4}$ -alkyl or  $-C(O)OR^{14}$  may be prepared by cyclisation of a compound of formula XXVIII A,



- wherein  $R^{12a}$  represents  $-S(O)_2-C_{1-4}$ -alkyl or  $-C(O)OR^{14}$  and n,  $R^6$ ,  $R^7$ ,  $R^{14}$  and  $L^2$  are as hereinbefore defined, for example at between  $0^\circ C$  and reflux temperature in the presence of a suitable base (e.g. sodium hydroxide), an appropriate solvent (e.g. dichloromethane, water, or a mixture thereof)

and, if necessary a phase transfer catalyst (such as tetrabutylammonium hydrogensulfate).

Compounds of formula VIII may be prepared by standard techniques. For example compounds of formula VIII in which:

(1) B represents  $-(CH_2)_nO-$  may be prepared by coupling a compound of formula XIA, as hereinbefore defined, to a compound of formula XXIX,



wherein  $L^4$  represents a suitable leaving group (e.g. halo) and n,  $R^5$ ,  $R^6$ , A and  $L^2$  are as hereinbefore defined; or

(2) B represents  $-C(O)N(R^{17})-$  may be prepared by coupling a compound of formula XXX,



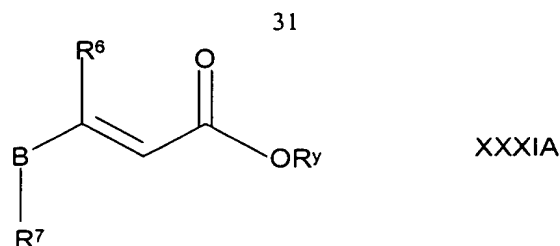
wherein  $R^7$  and  $R^{17}$  are as hereinbefore defined, to a compound of formula XXXI,



wherein  $L^4$ ,  $R^5$ ,  $R^6$ , A and  $L^2$  are as hereinbefore defined;

in both cases, under conditions which are well known to those skilled in the art.

Compounds of formula VIII in which A represents  $C_2$ -alkylene and  $R^5$  represents  $OR^{12}$ , in which  $R^{12}$  represents  $C_{1-6}$  alkyl or optionally substituted aryl may alternatively be prepared by reaction of a compound of formula XIX as hereinbefore defined with a compound of formula XXXIA,



wherein  $R^y$  represents  $C_{1-4}$  alkyl or aryl (which two groups are optionally substituted with one or more substituents selected from  $C_{1-4}$  alkyl or halo) and  $R^6$ ,  $R^7$  and B are as hereinbefore defined, for example at between ambient temperature (e.g.  $25^\circ\text{C}$ ) and reflux temperature in the presence of a suitable base (e.g.  $\text{K}_2\text{CO}_3$ ) and an appropriate organic solvent (e.g. acetonitrile), followed by conversion of the ester functionality to an  $L^2$  group (in which  $L^2$  is as hereinbefore defined), under conditions that are well known to those skilled in the art.

Compounds of formulae VII and VIII in which B represents  $-(\text{CH}_2)_n\text{S}(\text{O})-$  or  $-(\text{CH}_2)_n\text{S}(\text{O})_2-$  may be prepared by oxidation of corresponding compounds of formulae VII and VIII wherein B represents  $-(\text{CH}_2)_n\text{S}-$ , wherein n is as hereinbefore defined, in the presence of an appropriate amount of a suitable oxidising agent (e.g. *m*CPBA) and an appropriate organic solvent.

Compounds of formulae IX and XI may be prepared in a similar fashion to compounds of formula I (see, for example, process steps (a), (b), (c) or (d)).

Alternatively, compounds of formula IX in which A represents  $\text{C}_2$  alkylene may be prepared by reaction of a compound of formula VI, as hereinbefore defined with a compound of formula XXXII,



wherein B and  $R^7$  are as hereinbefore defined, for example a room temperature in the presence of a suitable organic solvent (e.g. ethanol).

Compounds of formula XIII may be prepared by removing an optionally substituted benzyloxycarbonyl unit from (i.e. deprotecting) a corresponding compound of formula I in which R<sup>7</sup> represents optionally substituted phenyl, R<sup>5</sup> and R<sup>6</sup> both represent H, B represents  
 5 -N(R<sup>17</sup>)C(O)O(CH<sub>2</sub>)-, A represents A<sup>a</sup> and A<sup>a</sup> is as hereinbefore defined under conditions which are well known to those skilled in the art.

Compounds of formula XV may be prepared by reaction of a  
 10 corresponding compound of formula I, as hereinbefore defined, in which R<sup>5</sup> represents -OH, with a compound of formula XXXIII



wherein R<sup>y</sup> is as hereinbefore defined, for example at between -10 and 25°C in the presence of a suitable solvent (e.g. dichloromethane),  
 15 followed by reaction with a suitable source of the azide ion (e.g. sodium azide) for example at between ambient and reflux temperature in the presence of an appropriate solvent (e.g. DMF) and a suitable base (e.g. NaHCO<sub>3</sub>).

20 Compounds of formula XV may alternatively be prepared by reaction of a corresponding compound of formula VI, as hereinbefore defined with a compound of formula XXXIIIA,



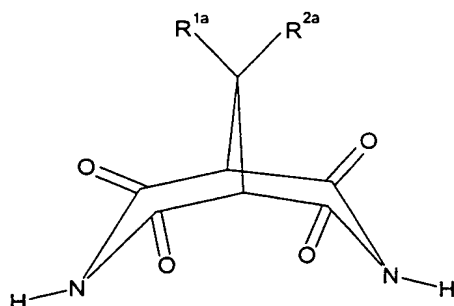
wherein L<sup>2</sup>, R<sup>6</sup>, R<sup>7</sup>, A and B are as hereinbefore defined, for example  
 25 under analogous conditions to those described hereinbefore for preparation of compounds of formula I (process step (e)).



Compounds of formula XX may be prepared by replacement of the OH group of a compound of formula I in which  $R^5$  represents OH with an  $L^2$  group under conditions that are well known to those skilled in the art.

- 5 Compounds of formula XXIA may be prepared by reaction of a corresponding compound of formula X with hydroxylamine, for example at elevated temperature (e.g. at reflux) in the presence of a suitable organic solvent (e.g. methanol).

- 10 Compounds of formula XXII are known in the literature or are readily available using known techniques. For example, compounds of formula XXII in which  $R^1$  and  $R^2$  together represent  $-O-(CH_2)_2-O-$ ,  $-(CH_2)_3-$ ,  $-(CH_2)_4-$  or  $-(CH_2)_5-$ , and  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$ ,  $R^{44}$ ,  $R^{45}$  and  $R^{46}$  all represent H, may be prepared by reduction of a compound of formula XXXIV,

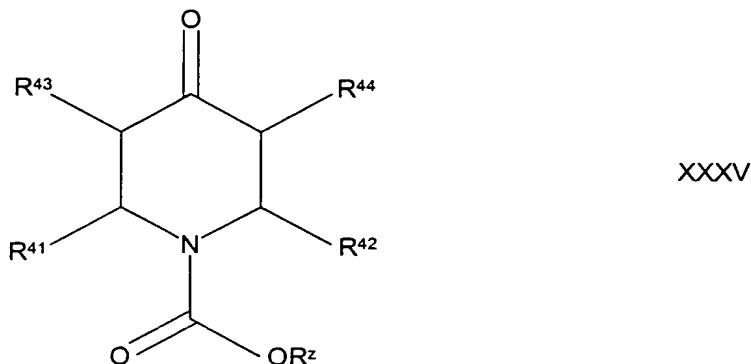


XXXIV

- 15 wherein  $R^{1a}$  and  $R^{2a}$  together represent  $-O-(CH_2)_2-O-$ ,  $-(CH_2)_3-$ ,  $-(CH_2)_4-$  or  $-(CH_2)_5-$ , in the presence of a suitable reducing agent (e.g.  $LiAlH_4$ ) under conditions which are well known to those skilled in the art.
- 20

Compounds of formula XXXIIIA may be prepared in analogous fashion to compounds of formula XV (i.e. from the corresponding alcohol).

Compounds of formulae X, XXIII and XXV (in which, in all cases,  $R^{45}$  and  $R^{46}$  both represent H), may be prepared, advantageously, by reaction of (as appropriate) either (i) a compound of formula XXXV,



- 5 wherein  $R^z$  represents  $C_{1-10}$  alkyl or  $C_{1-3}$  alkylaryl (e.g. alkylphenyl, such as benzyl) and  $R^{41}$ ,  $R^{42}$ ,  $R^{43}$  and  $R^{44}$  are as hereinbefore defined, or (ii) 4-piperidone (or a protected derivative thereof), with (as appropriate) either (1) a compound of formula XXXVI,



- 10 wherein  $R^5$ ,  $R^6$ ,  $R^7$ , A and B are as hereinbefore defined, or (2)  $NH_3$  (or a protected (e.g. benzyl) derivative thereof), in all cases in the presence of a formaldehyde (i.e. an appropriate source of formaldehyde, such as paraformaldehyde or formalin solution) and, in the case of compounds of formulae X and XXV, conversion of the  $C(O)OR^z$  group in the resultant  
 15 intermediate to a  $C(O)N(R^3)(R^4)$  group using techniques such as those described herein (e.g. process step (c) above).

The formation of compounds of formulae X, XXIII and XXV may be carried out in this way for example at between room temperature and reflux  
 20 (depending upon the concentration of the reactants) in the presence of an appropriate solvent (e.g. ethanol or methanol) and, preferably, in the presence of an organic acid (e.g. a  $C_{1-6}$  carboxylic acid, especially acetic acid).

It will be also appreciated by those skilled in the art that compounds of formula XXII in which  $R^1$  and  $R^2$  both represent H may also be prepared *via* this method (i.e. by reaction of a compound of 4-piperidone (or a  
 5 protected derivative thereof) with  $NH_3$  (or a protected derivative thereof) in the presence of a formaldehyde), provided that the intermediate so formed is subsequently reduced under appropriate reaction conditions.

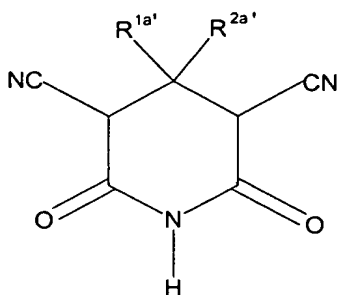
The skilled person will also appreciate that this process may also be used  
 10 to prepare compounds of formula I in which  $R^{41}$  and  $R^{42}$  are H, and  $R^{45}$  and/or  $R^{46}$  are other than H, for example by:

- (i) reacting a compound of formula XXXV in which  $R^{41}$  and/or  $R^{42}$  is/are other than H with, for example, benzylamine or a derivative thereof;
- 15 (ii) removal of the  $-C(O)OR^z$  unit;
- (iii) reaction at the free bispidine nitrogen of the resultant compound with a compound of formula VIII as hereinbefore defined;
- (iv) removal of the benzyl protecting group; and
- (v) reaction at the free bispidine nitrogen of the resultant compound  
 20 with, for example, a compound of formula III or IV as hereinbefore defined,

under conditions well known to those skilled in the art including those described hereinbefore. This reaction will be accompanied by, at some point, conversion of the bridgehead carbonyl functionality to the desired  
 25  $R^1/R^2$  groups.

Compounds of formula XXXIV may be prepared in accordance with techniques which are well known to those skilled in the art. For example, compounds of formula XXXIV in which  $R^{1a}$  and  $R^{2a}$  together represent

$-(CH_2)_3-$ ,  $-(CH_2)_4-$  or  $-(CH_2)_5-$  may be prepared by reaction of a compound of formula XXXVII,



XXXVII

wherein  $R^{1a'}$  and  $R^{2a'}$  together represent  $-(CH_2)_3-$ ,  $-(CH_2)_4-$  or  $-(CH_2)_5-$ ,  
 5 with a mixture of phosphoric acid and sulfuric acid, for example at  $120^\circ\text{C}$ .

Compounds of formula XXXVI are well known in the literature or are readily available using known techniques. For example, compounds of formula XXXVI wherein  $R^5$  represents OH,  $R^6$  represents H and A represents  $CH_2$  may be prepared by reaction of a compound of formula VII in which  $R^6$  represents H and X represents O with ammonium hydroxide under conditions which are well known to those skilled in the art.

15 Compounds of formulae III, VA, XIA, XII, XIV, XVI, XVII, XVIII, XIX, XXI, XXIB, XXIV, XXVI, XXVII, XXVIII, XXVIII A, XXIX, XXX, XXXI, XXXIA, XXXII, XXXIII, XXXV and XXXVII and derivatives thereof, are either commercially available, are known in the literature, or may be obtained either by analogy with the processes  
 20 described herein, or by conventional synthetic procedures, in accordance with standard techniques, from readily available starting materials using appropriate reagents and reaction conditions.

Substituents on the aryl (e.g. phenyl), and (if appropriate) heterocyclic,  
 25 group(s) in compounds defined herein may be converted to other claimed

substituents using techniques well known to those skilled in the art. For example, nitrobenzene may be reduced to an aminobenzene, hydroxy may be converted to alkoxy, alkoxy may be hydrolysed to hydroxy, etc.

- 5 The compounds of the invention may be isolated from their reaction mixtures using conventional techniques.

It will be appreciated by those skilled in the art that, in the process described above, the functional groups of intermediate compounds may be,  
10 or may need to be, protected by protecting groups.

Functional groups which it is desirable to protect include hydroxy, amino and carboxylic acid. Suitable protecting groups for hydroxy include trialkylsilyl and diarylalkylsilyl groups (e.g. *tert*-butyldimethylsilyl, *tert*-  
15 butyldiphenylsilyl or trimethylsilyl), tetrahydropyranyl and alkylcarbonyloxy groups (e.g. methyl- and ethylcarbonyloxy groups). Suitable protecting groups for amino include benzyl, *tert*-butyloxycarbonyl, 9-fluorenylmethoxycarbonyl or benzyloxycarbonyl. Suitable protecting groups for carboxylic acid include C<sub>1-6</sub> alkyl or benzyl esters.

20

The protection and deprotection of functional groups may take place before or after any of the reaction steps described hereinbefore.

Protecting groups may be removed in accordance with techniques which are  
25 well known to those skilled in the art and as described hereinafter.

The use of protecting groups is fully described in "Protective Groups in Organic Chemistry", edited by J W F McOmie, Plenum Press (1973), and

"Protective Groups in Organic Synthesis", 2nd edition, T W Greene & P G M Wutz, Wiley-Interscience (1991).

Persons skilled in the art will appreciate that, in order to obtain compounds of the invention in an alternative, and, on some occasions, more convenient, manner, the individual process steps mentioned herein may be performed in a different order, and/or the individual reactions may be performed at a different stage in the overall route (i.e. substituents may be added to and/or chemical transformations performed upon, different intermediates to those associated hereinbefore with a particular reaction). This will depend *inter alia* on factors such as the nature of other functional groups present in a particular substrate, the availability of key intermediates and the protecting group strategy (if any) to be adopted. Clearly, the type of chemistry involved will influence the choice of reagent that is used in the said synthetic steps, the need, and type, of protecting groups that are employed, and the sequence for accomplishing the synthesis.

It will also be appreciated by those skilled in the art that, although certain protected derivatives of compounds of formula I, which may be made prior to a final deprotection stage, may not possess pharmacological activity as such, they may be administered parenterally or orally and thereafter metabolised in the body to form compounds of the invention which are pharmacologically active. Such derivatives may therefore be described as "prodrugs". Moreover, we have found that certain compounds of formula I may act as prodrugs of other compounds of formula I.

All prodrugs of compounds of formula I are included within the scope of the invention.

Some of the intermediates referred to hereinbefore are novel. According to a further aspect of the invention there is thus provided: (a) a compound of formula II, as hereinbefore defined or a protected derivative thereof, provided that  $R^7$  does not represent optionally substituted phenyl; (b) a compound of formula V, as hereinbefore defined or a protected derivative thereof, provided that  $R^7$  does not represent optionally substituted phenyl; (c) a compound of formula X as hereinbefore defined or a protected derivative thereof; (d) a compound of formula XI as hereinbefore defined or a protected derivative thereof; (e) a compound of formula XIII, as hereinbefore defined or a protected derivative thereof; (f) a compound of formula XV, as hereinbefore defined or a protected derivative thereof; (g) a compound of formula XX, as hereinbefore defined or a protected derivative thereof; (h) a compound of formula XXIII, as hereinbefore defined or a protected derivative thereof, provided that  $R^7$  does not represent optionally substituted phenyl; and (i) a compound of formula XXV, as hereinbefore defined or a protected derivative thereof.

#### Medical and pharmaceutical use

20

The compounds of the invention are useful because they possess pharmacological activity. They are therefore indicated as pharmaceuticals.

Thus, according to a further aspect of the invention there is provided the compounds of the invention for use as pharmaceuticals.

In particular, the compounds of the invention exhibit myocardial electrophysiological activity, for example as demonstrated in the test described below.

- 5 The compounds of the invention are thus expected to be useful in both the prophylaxis and the treatment of arrhythmias, and in particular atrial and ventricular arrhythmias.

10 The compounds of the invention are thus indicated in the treatment or prophylaxis of cardiac diseases, or in indications related to cardiac diseases, in which arrhythmias are believed to play a major role, including ischaemic heart disease, sudden heart attack, myocardial infarction, heart failure, cardiac surgery and thromboembolic events.

- 15 In the treatment of arrhythmias, compounds of the invention have been found to selectively delay cardiac repolarization, thus prolonging the QT interval, and, in particular, to exhibit class III activity. Although compounds of the invention have been found to exhibit class III activity in particular, in the treatment of arrhythmias, their mode(s) of activity is/are  
20 not necessarily restricted to this class.

- According to a further aspect of the invention, there is provided a method of treatment of an arrhythmia which method comprises administration of a therapeutically effective amount of a compound of the invention to a person  
25 suffering from, or susceptible to, such a condition.



## Pharmaceutical preparations

The compounds of the invention will normally be administered orally, subcutaneously, intravenously, intraarterially, transdermally, intranasally, by inhalation, or by any other parenteral route, in the form of pharmaceutical preparations comprising the active ingredient either as a free base, a pharmaceutically acceptable ion exchanger or a non-toxic organic or inorganic acid addition salt, in a pharmaceutically acceptable dosage form. Depending upon the disorder and patient to be treated, as well as the route of administration, the compositions may be administered at varying doses.

The compounds of the invention may also be combined with any other drugs useful in the treatment of arrhythmias and/or other cardiovascular disorders.

According to a further aspect of the invention there is thus provided a pharmaceutical formulation including a compound of the invention in admixture with a pharmaceutically acceptable adjuvant, diluent or carrier.

Suitable daily doses of the compounds of the invention in therapeutic treatment of humans are about 0.05 to 5.0 mg/kg body weight at parenteral administration.

The compounds of the invention have the advantage that they are effective against cardiac arrhythmias.

Compounds of the invention may also have the advantage that they may be more efficacious than, be less toxic than, have a broader range of activity (including exhibiting any combination of class I, class II, class III and/or class IV activity (especially class I, class II and/or class IV activity in addition to class III activity)) than, be more potent than, be longer acting than, produce fewer side effects (including a lower incidence of proarrhythmias such as *torsades de pointes*) than, be more easily absorbed than, or that they may have other useful pharmacological properties over, compounds known in the prior art.

## Biological Tests

### Test A

#### Primary Electrophysiological Effects In Anaesthetised Guinea Pigs

Guinea pigs weighing between 660 and 1100 g were used. The animals were housed for at least one week before the experiment and had free access to food and tap water during that period.

Anaesthesia was induced by an intraperitoneal injection of pentobarbital (40 to 50 mg/kg) and catheters were introduced into one carotid artery (for blood pressure recording and blood sampling) and into one jugular vein (for drug infusions). Needle electrodes were placed on the limbs for recording of ECGs (lead II). A thermistor was placed in the rectum and the animal was placed on a heating pad, set to a rectal temperature of between 37.5 and 38.5°C.

A tracheotomy was performed and the animal was artificially ventilated with room air by use of a small animal ventilator, set to keep blood gases within

the normal range for the species. In order to reduce autonomic influences both vagi were cut in the neck, and 0.5 mg/kg of propranolol was given intravenously, 15 minutes before the start of the experiment.

- 5 The left ventricular epicardium was exposed by a left-sided thoracotomy, and a custom-designed suction electrode for recording of the monophasic action potential (MAP) was applied to the left ventricular free wall. The electrode was kept in position as long as an acceptable signal could be recorded, otherwise it was moved to a new position. A bipolar electrode  
10 for pacing was clipped to the left atrium. Pacing (2 ms duration, twice the diastolic threshold) was performed with a custom-made constant current stimulator. The heart was paced at a frequency just above the normal sinus rate during 1 minute every fifth minute throughout the study.
- 15 The blood pressure, the MAP signal and the lead II ECG were recorded on a Mingograph ink-jet recorder (Siemens-Elema, Sweden). All signals were collected (sampling frequency 1000 Hz) on a PC during the last 10 seconds of each pacing sequence and the last 10 seconds of the following minute of sinus rhythm. The signals were processed using a custom-made program  
20 developed for acquisition and analysis of physiological signals measured in experimental animals (see Axenborg and Hirsch, Comput. Methods Programs Biomed. 41, 55 (1993)).

The test procedure consisted of taking two basal control recordings, 5  
25 minutes apart, during both pacing and sinus rhythm. After the second control recording, the first dose of the test substance was infused in a volume of 0.2 mL into the jugular vein catheter for 30 seconds. Three minutes later, pacing was started and a new recording was made. Five minutes after the previous dose, the next dose of test substance was

administered. Six to ten consecutive doses were given during each experiment.

### Data analysis

5

Of the numerous variables measured in this analysis, three were selected as the most important for comparison and selection of active compounds. The three variables selected were the MAP duration at 75 percent repolarization during pacing, the atrio-ventricular (AV) conduction time (defined as the interval between the atrial pace pulse and the start of the ventricular MAP) during pacing, and the heart rate (defined as the RR interval during sinus rhythm). Systolic and diastolic blood pressure were measured in order to judge the haemodynamic status of the anaesthetised animal. Further, the ECG was checked for arrhythmias and/or morphological changes.

15

The mean of the two control recordings was set to zero and the effects recorded after consecutive doses of test substance were expressed as percentage changes from this value. By plotting these percentage values against the cumulative dose administered before each recording, it was possible to construct dose-response curves. In this way, each experiment generated three dose-response curves, one for MAP duration, one for AV-conduction time and one for the sinus frequency (RR interval). A mean curve of all experiments performed with a test substance was calculated, and potency values were derived from the mean curve. All dose-response curves in these experiments were constructed by linear connection of the data points obtained. The cumulative dose prolonging the MAP duration by 10% from the baseline was used as an index to assess the class III electrophysiological potency of the agent under investigation ( $D_{10}$ ).

20

25

## Test B

### Metabolic Stability of Test Compounds

5 An *in vitro* screen was set up to determine the metabolic stability of the compounds of the invention.

The hepatic S-9 fraction from dog, man, rabbit and rat with NADPH as co-factor was used. The assay conditions were as follows: S-9 (3 mg/mL), NADPH (0.83 mM), Tris-HCl buffer (50 mM) at pH 7.4 and 10  $\mu$ M of test  
10 compound.

The reaction was started by addition of test compound and terminated after 0, 1, 5, 15 and 30 minutes by raising the pH in the sample to above 10 (NaOH; 1 mM). After solvent extraction, the concentration of test  
15 compound was measured against an internal standard by LC (fluorescence/UV detection).

The percentage of test compound remaining after 30 minutes (and thus  $t_{1/2}$ ) were calculated and used as a measure for metabolic stability.

20

The invention is illustrated by way of the following examples.

## **Examples**

### 25 General Experimental Procedures

Mass spectra were recorded on a Finnigan MAT TSQ 700 triple quadrupole mass spectrometer equipped with an electrospray interface (FAB-MS) and VG Platform II mass spectrometer equipped with an electrospray interface (LC-MS), a Hewlett Packard model 6890 gas chromatograph connected to a

Hewlett-Packard model 5973A mass spectrometer *via* a Hewlett Packard HP-5-MS GC column, or a Shimadzu QP-5000 GC/mass spectrometer (CI, methane).  $^1\text{H}$  NMR and  $^{13}\text{C}$  NMR measurements were performed on a BRUKER ACP 300 and Varian UNITY plus 400 and 500 spectrometers, operating at  $^1\text{H}$  frequencies of 300, 400 and 500 MHz respectively, and at  $^{13}\text{C}$  frequencies of 75.5, 100.6 and 125.7 MHz respectively. Alternatively,  $^{13}\text{C}$  NMR measurements were performed on a BRUKER ACE 200 spectrometer at a frequency of 50.3 MHz.

- 10 Rotamers may or may not be denoted in spectra depending upon ease of interpretation of spectra. Unless otherwise stated, chemical shifts are given in ppm with the solvent as internal standard.

### Synthesis of intermediates

15

#### Example A

##### 4-(2-Oxiranylmethoxy)benzonitrile

- Epichlorohydrin (800 mL) and  $\text{K}_2\text{CO}_3$  (414 g) were added to a stirred solution of *p*-cyanophenol (238 g) in 2.0 L MeCN and the reaction mixture was refluxed under an inert atmosphere for 2 h. The hot solution was filtered and the filtrate concentrated, giving a clear oil which was crystallized from di-*iso*-propyl ether giving the product in 75% yield.

25

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  44.4, 49.7, 69.0, 104.5, 115.3, 118.9, 134.0, 161.6

#### Example B

##### 2(*S*)-Oxiranylmethyl 3-nitrobenzenesulfonate

*m*-Nitrobenzenesulfonylchloride (12.6 g; 57 mmol) was added to a cold

(-20°C) solution of (*R*)-(+)-glycidol (5.5 g; 74 mmol) and TEA (10.3 mL; 74 mmol). The reaction mixture was stirred at -20°C for 96 h. The solution was filtered and the filtrate washed with tartaric acid (10% w/w), brine, H<sub>2</sub>O and concentrated giving the title compound in a 97% yield.

5

<sup>1</sup>H NMR (CDCl<sub>3</sub>): δ 2.62 (dd, 1H), 2.84 (dd, 1H), 3.22 (m, 1H), 4.07 (dd, 1H), 4.49 (dd, 1H), 7.80 (t, 1H), 8.25 (m, 1H), 8.52 (m, 1H), 8.78 (m, 1H)

#### 10 Example C

##### 4-[(2*S*)-Oxiranylmethoxy]benzonitrile

The title compound was prepared in a 90% yield according to the procedure described in Example A above starting from (*R*)-(-)-epichlorohydrin.

15

#### Example D

##### 4-[(2*R*)-Oxiranylmethoxy]benzonitrile

The title compound was prepared according to the procedure described in Example A above starting from (*S*)-(-)-epichlorohydrin.

20

$[\alpha]_D^{20} = -14.1^\circ$  (*c* = 1.0; acetone)

<sup>1</sup>H NMR (CDCl<sub>3</sub>): δ 2.79 (1H, m); 2.98 (1H, m); 3.39 (1H, m); 3.98 (1H, m); 4.37 (1H, m); 6.99 (2H, d); 7.60 (2H, d)

25

Example E3-Benzyl-3,7-diazabicyclo[3.3.1]nonane(a) 3,7-Dibenzyl-3,7-diazabicyclo[3.3.1]nonane

- 5 The sub-title compound was prepared according to the method described in J. Org. Chem. **41**, 1593, (1976) except that 3,7-dibenzyl-3,7-diazabicyclo[3.3.1]nonan-9-one (also prepared according to the method described in J. Org. Chem. **41**, 1593 (1976)) was used instead of *N*-benzyl-*N*-methylbispidone.

(b) 3-Benzyl-3,7-diazabicyclo[3.3.1]nonane

- 3,7-Dibenzyl-3,7-diazabicyclo[3.3.1]nonane (1.97 g; 6.4 mmol; from step (a) above) was dissolved in EtOH (95%) and hydrogenated over 5% Pd/C at 1 atm. until tlc indicated that the reaction was complete. The catalyst was removed by filtration through a pad of Celite® and the residue was concentrated under reduced pressure to give the title compound in a quantitative yield.

- <sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 30.1, 33.4, 36.0, 52.5, 59.6, 64.3, 126.9, 128.3, 128.7, 138.8

Example F*tert*-Butyl 3,7-diazabicyclo[3.3.1]nonane-3-carboxylate

- 25 (a) *tert*-Butyl 7-benzyl-9-oxy-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate  
Paraformaldehyde (4.00 g; 127 mmol) was added to a solution of benzylamine (13.7 g; 126 mmol) in ethanol (190 mL). The solution was heated to 60°C and a solution of acetic acid (15.2 g; 252 mmol) in ethanol (160 mL) was added over 2 hours. After additional stirring for 1 hour, the



solution was cooled to room temperature. This solution was added (over 2 hours) to a mixture of 1-*tert*-butoxycarbonyl-4-piperidone (25.5 g; 127 mmol) and paraformaldehyde (4.80 g; 152 mmol) in ethanol (270 mL) which had been heated to 60°C. After reflux overnight, the solution was cooled to room temperature. The ethanol was removed by evaporation. Extractive work-up was performed in toluene:water and the material was filtered through silica in a toluene:ethyl acetate system. Evaporation of the eluant gave a solid material (37.4 g). The purity was 90 area% (HPLC) and the yield was 60%. By performing a crystallisation in *iso*-propanol, a compound with a purity of 98 area% (HPLC) and a yield of 70% was obtained.

MS (EI; 70 eV):  $m/z$  91 (100%),  $m/z$  57 (42%),  $m/z$  273 (32%),  $m/z$  330 (5%)

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  28.72, 47.71, 49.91, 50.60, 58.83, 59.16, 61.96, 80.18, 127.37, 128.45, 128.89, 137.57, 154.89, 213.66 (using TMS as reference)

(b) *tert*-Butyl 7-benzyl-9-oxy-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate  
(alternative preparation)

Benzylamine (6.51 g; 60.2 mmol), acetic acid (72.3 g, 1200 mmol), paraformaldehyde (3.71 g; 120 mmol) and 1-*tert*-butoxycarbonyl-4-piperidone (12.0 g; 60.2 mmol), were added to ethanol (300 mL). The solution was heated to 65°C and stirred at this temperature for 2 hours.

The same work-up procedure as that described in step (a) above was performed, yielding 15.78 g of material with a purity of 92 area% (HPLC) and a yield of 70%. Recrystallisation from *iso*-propanol yielded a compound with a purity of 94 area% (HPLC) in a yield of 54%.

(c) tert-Butyl 7-benzyl-3,7-diazabicyclo[3.3.1]-nonane-3-carboxylate

A mixture of 4-toluenesulfonehydrazide (12.4 mmol; 2.30 g) and *tert*-butyl 7-benzyl-9-oxy-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate (10.1 mmol; 4.00 g; 83.3%; from step (a) above) were dissolved in *iso*-propanol (30 mL) and heated at reflux for 2 hours. Acetic acid (2.5 mmol; 0.15 g) and sodium cyanoborohydride (12.1 mmol, 0.76 g) were added and the mixture was again heated at reflux for 2 hours. The slurry was cooled to ambient temperature and filtered. The filtrate was concentrated and an extractive work-up was performed in toluene:water. The toluene solution was concentrated to give 0.95 g of sub-title compound, with a purity of 90 area% (GC) in a yield of 60%.

MS (EI; 70 eV):  $m/z$  259 (100%),  $m/z$  91 (95%),  $m/z$  169 (45%),  $m/z$  57 (35%),  $m/z$  316 (25%)

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  28.67, 28.95, 31.11, 47.55, 48.38, 58.70, 58.96, 63.46, 78.71, 126.57, 128.00, 128.53, 138.94, 155.20 (using TMS as a reference)

(d) tert-Butyl 3,7-diazabicyclo[3.3.1]nonane-3-carboxylate

*tert*-Butyl 7-benzyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate (from step (c) above) was debenzylated according to the method described in Example E(b) above to give the title compound in quantitative yield.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  28.05, 28.29, 31.33, 48.35, 49.11, 51.53, 79.34, 155.16

Example G4-[3-(3,7-Diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile

HCl-saturated EtOAc (600 mL) was added to a solution of *tert*-butyl 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate (62 g; see Example 2 of international patent application No. PCT/SE98/02276) in EtOAc (600 mL) and the mixture was stirred at rt. for 4 h. The solvent was removed under reduced pressure, the residue was dissolved in MeCN (1.3 L) and K<sub>2</sub>CO<sub>3</sub> (100 g) was added. The suspension was stirred for 12 h and filtered. Concentration of the filtrate gave the title compound in a 90% yield.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 28.9, 29.2, 32.3, 50.9, 57.7, 60.8, 62.1, 66.0, 71.2, 104.0, 115.3, 119.1, 133.9, 162.1

(The title compound was also readily converted to the hydrochloride salt using standard techniques.)

Preparation of Compounds of Formula IExample 17-[(2S)-3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

Ethyl isocyanate (1.42 g, 16.6 mmol) was added to a solution of 4-{[(2S)-3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropyl]oxy}benzonitrile

(5.0 g, 20 mmol, see Example G above) in 30 mL of dichloromethane. The mixture was stirred for 4 hours at room temperature and was then concentrated *in vacuo* and purified by column chromatography on silica, eluting with dichloromethane: methanol (95:5), to yield 3.2 g (51%) of the title compound.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  15.52, 29.19, 29.50, 31.89, 35.77, 48.00, 49.17, 57.21, 60.49, 61.83, 65.41, 70.71, 103.88, 115.34, 119.15, 133.78, 133.84, 158.87, 162.19

## 5 Example 2

7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-(cyclopropylmethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

### (a) Cyclopropylmethyl isocyanate

10 Cyclopropylmethylamine (1.4 g, 19.7 mmol) was added to a suspension of 1,1'-carbonyldiimidazole (3.2 g, 19.7 mmol) in THF (10 mL). The resulting solution was stirred overnight at room temperature before being subjected to distillation, yielding 0.4 g (21 %) of the sub-title compound.

### 15 (b) 7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-(cyclopropylmethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

Cyclopropylmethyl isocyanate (0.4 g, 4 mmol, from step (a) above) was added to a solution of 4-[3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile (1.2 g, 4 mmol, see Example G above) in DCM.

20 The solution was stirred overnight, then concentrated *in vacuo*. The resulting residue was purified by column chromatography on silica gel, eluting with dichloromethane:methanol (93:7), to yield 0.85 g (50%) of the title compound.

25  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  3.29, 11.21, 29.31, 29.61, 32.10, 46.11, 48.14, 49.39, 57.24, 60.58, 62.04, 65.46, 70.76, 104.03, 115.37, 119.18, 133.88, 158.97, 162.22

Example 34-((2S)-2-Hydroxy-3-[7-(4-morpholinylcarbonyl)-3,7-diazabicyclo-[3.3.1]non-3-yl]propyl}oxy)benzonitrile

A solution of 4-[[2S)-3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxy-propyl]oxy}benzonitrile) (2.0 g, 6.6 mmol, prepared analogously to the method described in Example G above) in DCM (10 mL) was treated with aqueous NaOH (0.8 mL of 10 M), followed by 4-morpholinecarbonyl chloride (1.2 g, 8 mmol). The resulting mixture was stirred for 30 min. at room temperature, before water was added. The organic layer was separated, washed with 2 M NaOH followed by brine, before being separated, dried (MgSO<sub>4</sub>) and concentrated *in vacuo*. The residue was recrystallised twice, firstly from *iso*-propanol and then from ethanol, to yield 0.73 g (26.5%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 23.36, 29.59, 30.05, 32.34, 47.45, 49.51, 52.18, 56.86, 60.78, 62.82, 65.35, 66.66, 70.82, 104.03, 115.33, 119.17, 133.88, 162.23, 164.99

Example 47-{3-(4-Cyanophenoxy)-2-[(methanesulfonyl)amino]-propyl}-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide(a) 4-(3-Amino-2-hydroxypropoxy)benzonitrile

4-(2-Oxiranylmethoxy)benzonitrile (100 g, 0.57 mol, see Example A above) was added to a mixture of concentrated aqueous ammonium hydroxide (500 mL) and *iso*-propanol (300 mL). The resulting slurry was stirred at room temperature for 3 days. The reaction mixture was filtered to remove the insoluble by-product, and the filtrate was concentrated *in*

*vacuo* to give a crude product, which was crystallised from acetonitrile to yield 50 g (46%) of the sub-title compound.

(b) 2-(4-Cyanophenoxy)-1-{[(methanesulfonyl)amino]methyl}ethyl

5 methanesulfonate

Methanesulfonyl chloride (17.5 g, 153 mmol) was slowly added to a cooled (-10°C) solution of 4-(3-amino-2-hydroxypropoxy)benzonitrile (13.3 g, 69 mmol, from step (a) above) and 4-(dimethylamino)pyridine (0.2 g, 1.64 mmol) in pyridine (100 mL). The yellow solution was stirred  
10 at rt for 1.5 hours, concentrated *in vacuo* and then redissolved in DCM. This solution was washed twice with 2 M HCl and once with NaHCO<sub>3</sub> solution before the organic phase was separated, dried (MgSO<sub>4</sub>) and concentrated *in vacuo* to yield 23.5 g (100%) of the sub-title compound.

15 (c) 4-{[1-(Methanesulfonyl)aziridin-2-yl]methoxy}benzonitrile

A stirred solution of 2-(4-cyanophenoxy)-1-{[(methanesulfonyl)amino]-methyl}ethyl methanesulfonate (23.5 g, 67 mmol, from step (b) above) in acetonitrile (200 mL), was treated with potassium carbonate (30 g, 210 mmol), forming a thick precipitate. After 1 hour, a further portion of  
20 K<sub>2</sub>CO<sub>3</sub> (30 g, 210 mmol) was added. Stirring was continued for 2 h at rt before the reaction mixture was filtered and the filtrate concentrated *in vacuo*. The resulting oil (13 g) was crystallised from toluene to give 8 g (47%) of the sub-title compound.

25 mp 79-81°C

(d) *N*-{2-(7-Benzyl-3,7-diazabicyclo[3.3.1]non-3-yl)-1-[(4-cyanophenoxy)-methyl]ethyl}methanesulfonamide

A mixture of 3-benzyl-3,7-diazabicyclo[3.3.1]nonane (2 g, 10 mmol, see Example E above) and 4-[[1-(methanesulfonyl)aziridin-2-yl]methoxy]benzonitrile (2.5 g, 10 mmol, from step (c) above) in *iso*-propanol was refluxed overnight. The mixture was then concentrated *in vacuo*, giving a residue which was then dissolved in water (pH 3) and extracted with ether. The aqueous layer was made basic with 2 M NaOH and extracted with DCM. The dichloromethane layer was separated, dried and concentrated *in vacuo* to give a residue which was purified by column chromatography, eluting with a gradient of DCM:methanol:methanolic ammonia (98:2:0 to 97:0:3) to give 2.5 g (53%) of the sub-title compound.

(e) *N*-[2-(4-Cyanophenoxy)-1-(3,7-diazabicyclo[3.3.1]non-3-ylmethyl)-ethyl]methanesulfonamide

A solution of *N*-{2-(7-benzyl-3,7-diazabicyclo[3.3.1]non-3-yl)-1-[(4-cyanophenoxy)methyl]ethyl}methanesulfonamide (2.3 g 4.9 mmol, from step (d) above) in aqueous ethanol (95%; 55 mL) was hydrogenated over 5% Pd/C at ambient pressure. The catalyst was removed by filtration through a pad of Celite® and the residue was concentrated *in vacuo* to give 1.6 g of a crude product. This was recrystallised from methanol to yield 0.3 g (16%) of the sub-title compound.

(f) 7-{3-(4-Cyanophenoxy)-2-[(methanesulfonyl)amino]propyl}-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A suspension of *N*-[2-(4-cyanophenoxy)-1-(3,7-diazabicyclo[3.3.1]non-3-ylmethyl)ethyl]methanesulfonamide (0.29 g, 0.77 mmol, from step (e) above) in DCM (10 mL) was treated with ethyl isocyanate (66 µL,

0.84 mmol) to give a clear solution. The mixture was stirred for 1 h at rt, concentrated *in vacuo* and then purified by column chromatography, eluting with 5% MeOH in DCM, to give the title compound in 73% yield.

5  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  15.41, 28.88, 29.18, 30.77, 35.87, 41.78, 47.93, 48.65, 49.98, 58.24, 58.51, 60.15, 68.82, 104.51, 115.28, 118.95, 134.05, 158.58, 161.55

### Example 5

10 7-[(2*S*)-3-(4-Cyanophenoxy)-2-hydroxypropyl]-*N*-iso-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

(a) 7-Benzyl-*N*-iso-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

15 *iso*-Propyl isocyanate (1.7 g, 20 mmol) was slowly added to a solution of 3-benzyl-3,7-diazabicyclo[3.3.1]nonane (3.1 g, 14.3 mmol, see Example E above) in DCM (10 mL). The mixture was stirred at rt overnight and then concentrated *in vacuo* to yield 4.2 g (97%) of the sub-title compound.

(b) *N*-iso-Propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

20 A solution of 7-benzyl-*N*-iso-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (4.2 g, 14 mmol, from step (a) above) in methanol/water (17 mL of a 15:2 mixture) was hydrogenated over 5% Pd/C at ambient pressure. The catalyst was removed by filtration through a pad of Celite®, and the filtrate concentrated *in vacuo* to yield 2.6 g (87%) of the sub-title  
25 compound.



(c) 7-[(2*S*)-3-(4-Cyanophenoxy)-2-hydroxypropyl]-*N*-iso-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of 4-[(2*S*)-oxiranylmethoxy]benzonitrile (0.55 g, 3.14 mmol, see Example C above) and *N*-iso-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (0.85 g, 4 mmol, from step (b) above) in *iso*-propanol/water (6.5 mL of a 12:1 mixture) was stirred overnight at 60°C. The mixture was then concentrated *in vacuo* and the residue re-dissolved in DCM. The organic solution was washed with water then brine, dried (MgSO<sub>4</sub>) and concentrated *in vacuo* to give the title compound in 91 % yield.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 23.49, 29.29, 31.78, 42.26, 47.71, 49.09, 56.92, 60.27, 61.65, 65.19, 70.61, 103.54, 115.21, 119.09, 133.65, 158.11, 162.08

Example 6

7-[(2*R*)-3-(4-Cyano-2-[(2-cyanoethyl)amino]carbonyl)-phenoxy)-2-hydroxypropyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

(a) 7-Benzyl-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A cooled (0°C) solution of 3-benzyl-3,7-diazabicyclo[3.3.1]nonane (32.45 g, 0.15 mol, see Example E above) in DCM (300 mL) was treated with ethyl isocyanate (11.4 g, 0.16 mol), added dropwise. The solution was stirred for 2 h at rt before being concentrated *in vacuo*. The resulting residue was purified by chromatography on silica gel, eluting with a gradient of DCM:MeOH (100:0 to 90:10) to yield 36.4 g (84%) of the sub-title compound.

(b) N-Ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A solution of 7-benzyl-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (4.4 g, 15.3 mmol, from step (a) above) in aqueous ethanol (25 mL of 95%) was hydrogenated over 5% Pd/C at ambient pressure.

- 5 The catalyst was removed by filtration through a pad of Celite®, and the residue was concentrated *in vacuo* to yield 2.88 g (95%) of the sub-title compound.

(c) Methyl 5-bromo-2-hydroxybenzoate

- 10 Br<sub>2</sub> (52 g) was slowly added to a stirred solution of methyl salicylate (50 g; 330 mmol) in 300 mL acetic acid. The reaction mixture was stirred at rt. for 10 h, poured onto ice-water and the precipitate recrystallized from MeOH, giving the sub-title compound in a 83 % yield.

15 (d) Methyl 5-cyano-2-hydroxybenzoate

- Methyl 5-bromo-2-hydroxybenzoate (190.8 g; from step (c) above) and CuCN (73.9 g) were refluxed in DMF (500 mL) for 7 h. The temperature was allowed to decrease to 80°C and HCl (500 mL) and FeCl<sub>3</sub> (165.0 g) were added. The reaction mixture was stirred for 30 min., concentrated  
20 and partitioned between H<sub>2</sub>O and DCM. The organic layer was dried, concentrated the residue recrystallized from methylethyl ketone giving the sub-title compound in a 61 % yield.

(e) 5-Cyano-N-(2-cyanoethyl)-2-hydroxybenzamide

- 25 A mixture of methyl 5-cyano-2-hydroxybenzoate (20 g, 0.113 mol, from step (d) above), 3-aminopropanenitrile (15.4 g, 0.22 mol) and sodium cyanide (1 g, 20 mmol) in methanol (200 mL) was refluxed overnight. Tlc showed incomplete reaction, so DMSO (50 mL) was added, and reflux was continued for a further 5 h. The solution was concentrated *in vacuo*,

water added, followed by conc. HCl, until a precipitate formed. The product was filtered off, washed with water and dried to yield 19.4 g (80%) of the sub-title compound.

5 (f) 5-Cyano-*N*-(2-cyanoethyl)-2-[(2*R*)-oxiranylmethoxy]benzamide

A mixture of 5-cyano-*N*-(2-cyanoethyl)-2-hydroxybenzamide (2.1 g, 9.8 mmol, from step (e) above) and 10 equivalents of (S)-epichlorohydrin in *iso*-propanol:water (55 mL of 10:1) was refluxed overnight. The mixture was concentrated *in vacuo* and the residue purified by column chromatography, eluting with ethyl acetate to yield 0.63 g (24%) of the sub-title compound.

(g) 7-[(2*R*)-3-(4-Cyano-2-[(2-cyanoethyl)amino]carbonyl}phenoxy)-2-hydroxypropyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

15 A mixture of 5-cyano-*N*-(2-cyanoethyl)-2-[(2*R*)-oxiranylmethoxy]benzamide (0.63 g, 2.3 mmol, from step (f) above) and *N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (0.59 g, 3 mmol, from step (b) above) in *iso*-propanol:water (33 mL of 10:1) was stirred under reflux overnight. The reaction mixture was concentrated *in vacuo* and the residue purified by column chromatography, eluting with DCM:MeOH  
20 (9:1), to yield 0.78 g (73%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 15.40, 15.55, 17.94, 28.04, 29.21, 29.55, 31.31, 32.03, 35.69, 35.89, 36.21, 47.93, 48.65, 49.36, 57.00, 60.47, 61.05,  
25 65.32, 72.21, 105.39, 114.37, 118.22, 118.45, 123.28, 136.36, 136.45, 158.53, 159.20, 160.08, 163.75

ES-MS (M+1)<sup>+</sup> 469.0 (m/z)

Example 7

7-((2S)-3-{4-Cyano-2-[(cyclopropylamino)carbonyl]-phenoxy}-2-hydroxypropyl)-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

5

(a) N<sup>1</sup>-Cyclopropyl-5-cyano-2-hydroxybenzamide

Cyclopropylamine (14.3 g) and Na (100 mg) were added to a solution of methyl 5-cyano-2-hydroxybenzoate (10.0 g; from step (d) above) in DMSO (40 mL). The reaction mixture was heated at 80°C in a sealed steel vessel overnight, diluted with H<sub>2</sub>O, acidified and extracted with EtOAc, giving the sub-title compound (11.0 g), after concentration of the organic layer.

(b) 5-Cyano-N-cyclopropyl-2-[(2S)-oxiranylmethoxy]benzamide

A mixture of N<sup>1</sup>-cyclopropyl-5-cyano-2-hydroxybenzamide (1.56 g, 7.7 mmol, from step (a) above), (2S)-oxiranylmethyl 3-nitrobenzenesulfonate (2 g, 7.7 mmol, see Example B above) and K<sub>2</sub>CO<sub>3</sub> (1.16 g, 8.4 mmol) in 2-butanone (15 mL) was stirred at 60°C for 18 h. The mixture was concentrated *in vacuo* and the residue crystallised from di-*iso*-propyl ether:MeCN (9:1) to yield 0.97 g (97%) of the sub-title compound.

(c) 7-((2S)-3-{4-Cyano-2-[(cyclopropylamino)carbonyl]phenoxy}-2-hydroxypropyl)-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of 5-cyano-N-cyclopropyl-2-[(2S)-oxiranylmethoxy]benzamide (0.97 g, 3.8 mmol, from step (b) above) and N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (0.89 g, 4.5 mmol, see Example 6(b) above) in *iso*-propanol:water (22 mL of 10:1) was refluxed overnight. The solvent was removed *in vacuo* and the resulting residue purified by

column chromatography on silica gel, eluting with DCM:MeOH (9:1), to yield 1.37 g (79%) of the title compound.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  6.62, 6.78, 15.81, 23.55, 29.61, 29.90, 32.48, 36.20, 48.32, 49.84, 53.68, 57.48, 60.92, 62.06, 65.61, 71.72, 105.42, 113.69, 118.64, 123.78, 136.26, 136.77, 159.70, 159.97, 164.75

### Example 8

#### N-Ethyl-7-(4-nitrophenethyl)-3,7-diazabicyclo[3.3.1]nonane-3-

#### carboxamide

A mixture of 1-(2-bromoethyl)-4-nitrobenzene (1.6 g, 7.0 mmol), N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (1.0 g, 5.1 mmol, see Example 6(b) above) and  $\text{K}_2\text{CO}_3$  (1.38 g, 10 mmol) was stirred at rt overnight. The mixture was then filtered and concentrated *in vacuo* and the resulting residue purified by column chromatography, eluting with a gradient of DCM:MeOH (100:0 to 90:10), to yield 1.5 g (85%) of the title compound.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  15.71, 28.83, 30.11, 33.03, 35.67, 47.97, 59.22, 59.49, 123.34, 129.65, 146.26, 149.15, 157.95

### Example 9

#### N-(Cyanomethyl)-7-[(2S)-3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

#### (a) Cyanomethyl isocyanate

The title compound was prepared according to the procedure described in Example 2(a) above, using 2-aminoacetonitrile in place of cyclopropylmethylamine.

(b) N-(Cyanomethyl)-7-[(2S)-3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

The title compound was prepared in 26% yield (counting steps (a) and (b) together) according to procedure described in Example 2(b) above, using cyanomethyl isocyanate (from step (a) above) in place of cyclopropylmethyl isocyanate.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 28.99, 29.27, 29.47, 31.77, 48.32, 49.33, 56.88, 60.33, 61.61, 65.32, 70.63, 103.96, 115.31, 117.63, 119.21, 133.93, 157.74, 162.08

Example 10

N-Ethyl-7-{4-[(methanesulfonyl)amino]phenethyl}-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

(a) 4-[(Methanesulfonyl)amino]phenethyl methanesulfonate

Methanesulfonyl chloride (45 g, 0.39 mol) was added, dropwise over 30 minutes, to a cooled (-5°C) solution of 4-aminophenethyl alcohol (25.2 g, 0.18 mol) in pyridine (200 mL). The mixture was stirred at 0°C for 1 h and then at rt overnight. The resulting red suspension was poured in to a mixture of ice (300 mL) and conc. HCl (60 mL). The pink precipitate that formed was filtered off, redissolved in DCM, dried and treated with activated carbon. The resulting solution was concentrated *in vacuo* to give a residue, which, on recrystallisation from ethyl acetate, gave 34.5 g (64%) of the sub-title compound.

mp 133-134°C

(b) *N*-Ethyl-7-{4-[(methanesulfonyl)amino]phenethyl}-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide

A mixture of *N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (1 g, 5 mmol, see Example 6(b) above), 4-[(methanesulfonyl)amino]phenethyl methanesulfonate (1.5 g, 5 mmol, from step (a) above) and NaHCO<sub>3</sub> (3 g, 35.7 mmol) in MeCN (50 mL) was refluxed for 3 h under nitrogen. The reaction mixture was filtered and concentrated *in vacuo* to give 2.2 g of crude product, which was filtered through a silica plug, with MeOH/2 *N* HCl. The pH of the fractions was raised to pH 6 and extracted with DCM, yielding 0.2 g of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 15.75, 28.87, 30.23, 32.58, 35.64, 35.76, 39.14, 48.18, 59.17, 60.26, 121.41, 129.85, 134.72

15 Example 11

7-[3-(4-Cyanophenoxy)-2-fluoropropyl]-*N*-ethyl-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide

20 (a) 7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-*N*-ethyl-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide

The title compound was prepared according to the procedure described in Example 1 above, using 4-[3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile (see Example G above) in place of 4-[(2*S*)-3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropyl]oxy}benzonitrile.

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(b) 7-[3-(4-Cyanophenoxy)-2-fluoropropyl]-*N*-ethyl-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide

A solution of 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (1.0 g, 2.7 mmol, from step (a)

above) in DCM (2.5 mL) was cooled to -78°C. A solution of (diethylamino)sulfurtrifluoride in DCM (2.5 mL) was added slowly under stirring. Stirring was continued for 35 minutes, during which time the reaction was allowed to warm to room temperature. Dichloromethane was added and the reaction mixture was then washed with NaHCO<sub>3</sub>, dried and concentrated *in vacuo*. The resulting residue was purified by column chromatography, eluting with DCM:MeOH (98:2), to yield 0.68 g (67%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 15.63, 29.00, 30.33, 35.70, 47.78, 47.93, 58.36, 58.67, 59.82, 60.39, 68.60, 68.89, 89.56, 91.86, 104.15, 115.56, 119.25, 133.97, 157.61, 161.92

#### Example 12

7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-[2-oxo-2-(propylamino)-ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

(a) Ethyl 2-[(7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]non-3-yl}carbonyl)amino]acetate

A cooled (0°C) solution of 4-[3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile (23.1 g, 77 mmol, see Example G above) in DCM (700 mL) was treated with ethyl 2-isocyanatoacetate (9.92 g, 77 mmol), and then stirred at rt for 7 h. The reaction mixture was concentrated *in vacuo* to yield 33.6 g (100%) of the sub-title compound.

(b) 7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-[2-oxo-2-(propylamino)-ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of ethyl 2-[(7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]non-3-yl}carbonyl)amino]acetate (0.76 g 1.8 mmol,



from step (a) above), propylamine (5 mL, 3.6 g, 69.1 mmol) and NaCN (0.01 g, 0.2 mmol) in methanol (10 mL) was warmed to 75°C in a sealed tube overnight. The solvent was then removed *in vacuo* and the residue diluted with Na<sub>2</sub>CO<sub>3</sub> solution. The aqueous mixture was extracted with DCM, and the resulting organic layer separated, dried and concentrated *in vacuo*. The resulting residue was purified by column chromatography, eluting with a gradient of dichloromethane:methanol (100:0 to 90:10), to give the title compound in 70% yield.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 11.36, 22.65, 29.12, 29.42, 31.78, 41.15, 44.75, 48.15, 49.10, 56.99, 60.40, 61.35, 65.33, 70.74, 103.99, 115.27, 119.12, 133.91, 158.71, 162.10, 170.62

### Example 13

7-{3-(4-Cyanophenoxy)-2-[(4-morpholinylcarbonyl)amino]propyl}-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

#### (a) tert-Butyl 3-(4-cyanophenoxy)-2-hydroxypropylcarbamate

A cooled (0°C) solution of 4-(3-amino-2-hydroxypropoxy)benzonitrile (44.6 g, 0.23 mol, see Example 4(a) above) in THF:H<sub>2</sub>O (1.5 L of 1:1) was treated with di-*tert*-butyl dicarbonate (53 g, 0.24 mol). The mixture was stirred at rt overnight, after which NaCl was added and the resulting organic layer separated. The water layer was extracted with ether and the combined organics were dried and concentrated *in vacuo*. The resulting oil (70 g) was filtered through a plug of silica, and then crystallised from diethyl ether:di-*iso*-propyl ether to yield 50 g of the sub-title compound.

(b) 2-[(*tert*-Butoxycarbonyl)amino]-1-[(4-cyanophenoxy)methyl]ethyl methanesulfonate

Methanesulfonyl chloride (22.3 g 0.195 mol) was added over the course of 1.5 hours to a cooled (0°C) solution of *tert*-butyl 3-(4-cyanophenoxy)-2-hydroxypropylcarbamate (51.2 g, 0.177 mol, from step (a) above) and 4-(dimethylamino)pyridine (1.3 g, 10.6 mmol) in pyridine (250 mL), kept under an inert atmosphere. The reaction mixture was stirred for 2 h at rt before water and DCM were added. The organic layer was separated, washed with water, dried (MgSO<sub>4</sub>) and concentrated *in vacuo* to yield 68.1 g (100%) of the sub-title compound.

(c) *tert*-Butyl 2-[(4-cyanophenoxy)methyl]-1-aziridinecarboxylate

A cooled (0°C) solution of 2-[(*tert*-butoxycarbonyl)amino]-1-[(4-cyanophenoxy)methyl]ethyl methanesulfonate (30.6 g, 82.6 mmol, from step (b) above) and tetrabutylammonium hydrogensulfate (3 g, 8.8 mmol) in DCM (100 mL) was treated with 50 wt.% aqueous NaOH (60 mL) under an inert atmosphere. The resulting mixture was stirred, and the temperature was slowly allowed to rise to rt over for 4 h, and then extracted with ether. The organic layer was washed with water and concentrated *in vacuo* to give a residue that was purified by column chromatography (dichloromethane eluant). Crystallisation from diethyl ether:*di-iso*-propyl ether gave the sub-title compound in quantitative yield.

(d) *tert*-Butyl 2-(4-cyanophenoxy)-1-({7-[(ethylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl)methyl}ethyl)carbamate

A mixture of *N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (2.88 g, 14.6 mmol, see Example 6(b) above) and *tert*-butyl 2-[(4-cyanophenoxy)methyl]-1-aziridinecarboxylate (4.0 g, 14.6 mmol, from step (c) above) in *iso*-propanol (20 mL) was refluxed overnight. The

reaction mixture was concentrated *in vacuo* to give 7.4 g of a yellow oil, which was purified by column chromatography, eluting with a gradient of DCM:MeOH (100:0 to 90:10), to yield 3.33 g of the sub-title compound.

5 (e) 7-[2-Amino-3-(4-cyanophenoxy)propyl]-N-ethyl-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide

A solution of *tert*-butyl 2-(4-cyanophenoxy)-1-({7-[(ethylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl}methyl)ethylcarbamate (2.4 g, 5.1 mmol, from step (d) above) in HCl-saturated ethyl acetate was stirred for 1 h at  
10 rt. The reaction mixture was then concentrated *in vacuo* and resulting residue re-dissolved in water. The aqueous solution was treated with aqueous NaHCO<sub>3</sub> and extracted with DCM, which organic layer was then dried and concentrated *in vacuo* to give 2 g of the sub-title compound.

15 (f) 7-{3-(4-Cyanophenoxy)-2-[(4-morpholinylcarbonyl)amino]propyl}-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A cooled (5°C) solution of 7-[2-amino-3-(4-cyanophenoxy)propyl]-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (0.33 g, 0.7 mmol, from step (e) above) and triethylamine (0.4 mL, 3.0 mmol) in DCM  
20 (5 mL) was treated with 4-morpholinecarbonyl chloride (0.11 g, 0.7 mmol), and then stirred at 5°C for 3 h. After further stirring at room temperature overnight, tlc analysis indicated incomplete reaction, and so a further portion of 4-morpholinecarbonyl chloride (40 mg, 0.27 mmol) was added. Stirring was continued at rt overnight again before NaHCO<sub>3</sub>  
25 solution was added. The organic layer was separated, dried and concentrated *in vacuo* to give 400 mg of crude product, which was purified by column chromatography on silica gel, eluting with dichloromethane:methanolic ammonia (95:5) to give 250 mg of the title compound.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  161.94, 158.26, 157.81, 133.94, 119.15, 115.37, 103.90, 67.26, 66.66, 60.66, 60.51, 57.99, 48.93, 48.37, 47.39, 44.06, 35.93, 30.71, 29.34, 29.02, 15.51

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#### Example 14

#### N-(4-Cyanophenethyl)-7-(4-oxoheptyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

#### 10 (a) 3-Benzyl-7-[3-(2-propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]nonane

A mixture of 3-benzyl-3,7-diazabicyclo[3.3.1]nonane (10.5 g, 48.5 mmol, see Example E above), 2-(3-bromopropyl)-2-propyl-1,3-dioxolane (11.5 g, 48.5 mmol, Bajrowicz *et al.*, *Tetrahedron*, **41** (1985) 1833) and  $\text{K}_2\text{CO}_3$  (13.8 g, 0.1 mol) in MeCN (50 mL) was refluxed overnight. The reaction mixture was filtered and concentrated *in vacuo* to yield 18.8 g (100%) of the sub-title compound.

#### 20 (b) 3-[3-(2-Propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]-nonane

A solution of 3-benzyl-7-[3-(2-propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]nonane (18.8 g, 4.85 mmol, from step (a) above) in ethanol (100 mL) was hydrogenated over 5% Pd/C at ambient pressure. The catalyst was removed by filtration through a pad of Celite®, and the filtrate concentrated *in vacuo* to yield 13.7 g (100%) of the sub-title compound.

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(c) N-(4-Cyanophenethyl)-7-(4-oxoheptyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A solution of 4-(2-aminoethyl)benzonitrile (1.0 g, 6.9 mmol, Wiley *et al.*, *Bioorg. Med. Chem. Lett.*, 6 (1996) 2387) in dry THF (10 mL) was treated with 1,1'-carbonyldiimidazole (1.17 g, 7.2 mmol), and the mixture was stirred for 30 min. A solution of 3-[3-(2-propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]nonane (1.3 g, 4.6 mmol, from step (b) above) in THF (5 mL) was added to the reaction mixture, and stirring was continued overnight at rt. The solution was then concentrated *in vacuo* and the resulting residue diluted with MeOH and 2 M HCl, which solution was stirred for 2 h at rt. The mixture was made alkaline and extracted with DCM. The organic layer was separated, dried and concentrated *in vacuo* to give a residue which was purified by flash chromatography, eluting with DCM:MeOH (92:8), to yield 0.57 g (30%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 13.73, 17.21, 20.85, 28.79, 30.38, 36.91, 39.84, 41.83, 44.73, 47.94, 57.65, 59.05, 110.06, 118.93, 129.67, 132.20, 145.52, 157.47, 211.67

Example 15

N'-(4-Cyanobenzoyl)-7-(4-oxoheptyl)-3,7-diazabicyclo[3.3.1]nonane-3-carbohydrazide

A mixture of 4-cyanobenzohydrazide (0.82 g, 5.0 mmol) and 1,1'-carbonyldiimidazole (0.82 g, 5 mmol) in THF (15 mL) was stirred for 10 min at rt before 3-[3-(2-propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]nonane (1.44 g, 5.0 mmol, see Example 14(b) above) was added. The reaction mixture was stirred overnight at rt, before being concentrated *in vacuo*. The resulting residue was dissolved in DCM, and washed with water. The organic layer was separated and concentrated *in vacuo* to give

a residue which was dissolved in methanol/2M HCl. Evaporation of the MeOH *in vacuo* and extraction of the remaining aqueous solution with DCM, gave, after purification by flash chromatography on silica gel (dichloromethane:methanolic ammonia eluant), 0.5 g (25%) of the title compound.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  213.21, 164.24, 157.01, 136.31, 132.19, 128.24, 118.11, 115.11, 58.65, 57.89, 48.38, 44.31, 40.55, 31.52, 29.12, 21.60, 17.08, 13.69

#### Example 16

4-{2-Amino-3-[7-(1-piperidinylcarbonyl)-3,7-diazabicyclo[3.3.1]non-3-yl]propoxy}benzonitrile

(a) 7-Benzyl-3,7-diazabicyclo[3.3.1]non-3-yl(1-piperidinyl)methanone

The sub-title compound was prepared by way of a reaction between 3-benzyl-3,7-diazabicyclo[3.3.1]nonane (see Example E above) and 1-piperidinecarbonyl chloride (Boon, *J. Chem. Soc.*, (1947) **307**, 313).

(b) 3,7-Diazabicyclo[3.3.1]non-3-yl(1-piperidinyl)methanone

The sub-title compound was obtained in quantitative yield according to the procedure described in Example 14(b) above, using 7-benzyl-3,7-diazabicyclo[3.3.1]non-3-yl(1-piperidinyl)methanone (from step (a) above) in place of 3-benzyl-7-[3-(2-propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]nonane.

(c) *tert*-Butyl 2-(4-cyanophenoxy)-1-{[7-(1-piperidinylcarbonyl)-3,7-diaza-bicyclo[3.3.1]non-3-yl]methyl}ethylcarbamate

A mixture of *tert*-butyl 2-[(4-cyanophenoxy)methyl]-1-aziridinecarboxylate (1.92 g, 7 mmol, see Example 13(c) above) and 3,7-diaza-bicyclo[3.3.1]non-3-yl(1-piperidinyl)methanone (1.85 g, 7 mmol, from step (a) above) in *iso*-propanol (15 mL) was refluxed for 30 h. The solution was concentrated *in vacuo* to yield 3.7 g of crude product, which was purified by chromatography using 2.5% MeOH in DCM to give 2.0 g (56%) of sub-title compound.

(d) 4-{2-Amino-3-[7-(1-piperidinylcarbonyl)-3,7-diazabicyclo[3.3.1]non-3-yl]propoxy}benzonitrile

A cooled (0°C) solution of *tert*-butyl 2-(4-cyanophenoxy)-1-{[7-(1-piperidinylcarbonyl)-3,7-diazabicyclo[3.3.1]non-3-yl]methyl}ethyl-

carbamate (1.9 g, 3.7 mmol, from step (c) above) in ethyl acetate was treated with HCl-saturated ethyl acetate. The mixture was stirred for 4 h before being concentrated *in vacuo*. The resulting residue was dissolved in water, made basic with NaHCO<sub>3</sub> and extracted with DCM. The organic layer was separated, dried and concentrated *in vacuo* to yield 1.5 g (100%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 24.73, 25.72, 29.62, 29.95, 32.11, 47.44, 48.14, 49.53, 50.98, 57.87, 60.57, 62.59, 72.03, 103.90, 115.30, 119.22, 133.91, 162.23, 164.35

Example 17N-Ethyl-7-{2-hydroxy-3-[4-(1*H*-imidazol-1-yl)phenoxy]propyl}-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide5 (a) 1-[4-(2-Oxiranylmethoxy)phenyl]-1*H*-imidazole

A mixture of 4-(1*H*-imidazol-1-yl)phenol (10 g, 60 mmol), K<sub>2</sub>CO<sub>3</sub> (8.63 g, 60 mmol) and 2-oxiranylmethyl 3-nitrobenzenesulfonate (15.5 g, 60 mmol, see Example B above) in DMF (140 mL) was stirred at 40°C overnight. The mixture was then concentrated *in vacuo* and the resulting  
 10 residue diluted with DCM, washed with water, dried and then concentrated *in vacuo*. The crude product was then purified by flash chromatography, eluting with a gradient of dichloromethane:methanol (100:0 to 70:30) to yield 3.4 g, (72.6%) of the title compound.

15 (b) N-Ethyl-7-{2-hydroxy-3-[4-(1*H*-imidazol-1-yl)phenoxy]propyl}-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of 1-[4-(2-oxiranylmethoxy)phenyl]-1*H*-imidazole (3.16 g, 14.6 mmol, from step (a) above) and *N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (2.88 g 14.6 mmol, see Example 6(b)  
 20 above) in *iso*-propanol:H<sub>2</sub>O (18 mL of 9:1) was refluxed for 3 hours, concentrated *in vacuo* and purified by acid/base extraction to yield 4.4 g (72.6%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 15.52, 29.13, 29.44, 31.84, 35.70, 47.92, 49.07,  
 25 57.21, 60.44, 61.94, 65.45, 70.76, 115.49, 118.58, 122.90, 129.86, 130.56, 135.66, 158.16, 158.78



Example 18N-[3-(4-Cyanophenoxy)propyl]-7-(2-hydroxyethyl)-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide5 (a) 4-(3-Bromopropoxy)benzonitrile

1,3-Dibromopropane (1.02 L; 10 mol) was added to a stirred suspension of *p*-cyanophenol (238 g; 2 mol), K<sub>2</sub>CO<sub>3</sub> (276.4 g; 2 mol) in MeCN (2.7 L). The reaction mixture was refluxed for 4 h, filtered and concentrated. The residue was recrystallized from *iso*-propyl ether to give the sub-title  
10 compound in a 69% yield.

(b) 4-[3-(1,3-Dioxo-1,3-dihydro-2*H*-isoindol-2-yl)propoxy]benzonitrile

A mixture of 4-(3-bromopropoxy)benzonitrile (20 g, 84 mmol, see step (a) above) and potassium phthalimide (15.5 g, 84 mmol) in DMF (120 mL)  
15 was stirred at 95°C for 4 h. The solution was then concentrated *in vacuo* and the resulting residue dissolved in DCM and washed with water. The organic layer was separated, dried (Na<sub>2</sub>SO<sub>4</sub>) and concentrated *in vacuo* to yield 25.5 g (99%) of the sub-title compound.

20 (c) 4-(3-Aminopropoxy)benzonitrile

A mixture of 4-[3-(1,3-dioxo-1,3-dihydro-2*H*-isoindol-2-yl)propoxy]-benzonitrile (25.5 g, 83 mmol, from step (b) above) and hydrazine hydrate (4.15 g, 83 mmol) in methanol (100 mL) was refluxed for 1 h before water (120 mL) was added. The methanol was evaporated under reduced  
25 pressure and concentrated hydrochloric acid (120 mL) was added. The resulting mixture was heated on a steam bath for 1.5 h and then cooled in the refrigerator overnight. The resulting precipitate was filtered off and the filtrate was concentrated *in vacuo*. Water was added to the resulting residue and the solution made basic. The aqueous solution was extracted

with DCM, which organic layer was then separated, dried and concentrated *in vacuo* to yield 6 g (41 %) of the sub-title compound.

(d) 7-Benzyl-3,7-diazabicyclo[3.3.1]nonane-3-ethanol

- 5 The compound was prepared in 72% yield by reacting 3-benzyl-3,7-diazabicyclo[3.3.1]nonane (see Example E above) with 2-bromoethanol.

(e) 3,7-Diazabicyclo[3.3.1]nonane-3-ethanol

- 10 The sub-title compound was prepared according to the procedure described in Example 14(b) above, using 7-benzyl-3,7-diazabicyclo[3.3.1]nonane-3-ethanol (from step (d) above) in place of 3-benzyl-7-[3-(2-propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]nonane.

- 15 (f) N-[3-(4-Cyanophenoxy)propyl]-7-(2-hydroxyethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

- The title compound was prepared in 11 % yield according to the procedure described in Example 14(c) above, using 3,7-diazabicyclo[3.3.1]nonane-3-ethanol (from step (e) above) and 4-(3-aminopropoxy)benzonitrile (from step (c) above) in place of 3-[3-(2-propyl-1,3-dioxolan-2-yl)propyl]-3,7-diazabicyclo[3.3.1]nonane and 4-(2-aminoethyl)benzonitrile, respectively.
- 20

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 162.04, 158.99, 133.66, 118.99, 115.03, 103.35, 66.55, 60.24, 57.87, 57.18, 50.02, 48.63, 37.93, 31.81, 29.26, 28.96

Example 19N-{[7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]non-3-yl]carbonyl}-4-methylbenzenesulfonamide

A solution of 4-[3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile (200 mg, 0.66 mmol, see Example G above) in chloroform (20 mL) was treated with a solution of *p*-toluenesulfonyl isocyanate (110  $\mu$ L of 96% purity, 0.136 g, 0.69 mmol in chloroform (4 mL), added dropwise. A white precipitate immediately formed and the mixture was then concentrated *in vacuo*. The crude product so obtained was subjected to chromatography on silica gel, eluting with hexane:ethyl acetate:methanolic ammonia (75:75:50) to give the title compound in 53% yield.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  15.77, 29.18, 32.37, 36.13, 48.72, 52.27, 56.32, 109.83, 113.13, 118.27, 118.93, 120.10, 127.80, 131.39, 132.46, 132.73, 134.62, 138.75, 159.14, 167.09

Example 20N-Allyl-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of allylamine (125  $\mu$ L, 1.66 mmol) and 1,1'-carbonyl-diimidazole (269 mg, 1.66 mmol) in THF (10 mL) was stirred at rt for 40 min. The mixture was then treated with a solution of 4-[3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile (see Example G above) in THF (5 mL), and stirring continued overnight. The mixture was concentrated *in vacuo* and the resulting residue purified by chromatography on silica gel, eluting with hexane:methanolic ammonia (1:1) to give the title compound in 57% yield.

<sup>13</sup>C NMR (MeOD): δ 29.37, 30.79, 41.95, 42.91, 58.91, 59.55, 61.12, 66.52, 70.75, 103.31, 113.81, 115.39, 118.72, 133.73, 135.57, 136.06, 158.93, 162.67

5 Example 21

7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-[2-(2-thienyl)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

The title compound was prepared in 83% yield according to the procedure described in Example 19 above, using 2-(2-isocyanatoethyl)thiophene in place of *p*-toluenesulfonyl isocyanate.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 29.19, 29.50, 30.59, 32.11, 42.26, 47.94, 49.37, 56.23, 60.47, 61.95, 65.32, 70.74, 103.88, 115.36, 119.52, 123.69, 125.25, 127.04, 133.90, 142.19, 158.74, 162.22

15 Example 22

7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-[3-(ethylamino)-3-oxopropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

20 (a) Ethyl 3-[(7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]non-3-yl)carbonyl]amino]propanoate

The sub-title compound was prepared in 90% yield according to the procedure described in Example 12(a) above, using ethyl 3-isocyanatopropanoate in place of ethyl 2-isocyanatoacetate.

25 (b) 7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-[3-(ethylamino)-3-oxopropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

The title compound was prepared in 22% yield according to the procedure described in Example 12(b) above, using ethyl 3-[(7-[3-(4-

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*N*-(1-Cyanoethyl)-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

## 15

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A mixture of 2-aminopropanenitrile (250 mg, 3.58 mmol, from step (a) above) and *N*-ethyl di-*iso*-propylamine (0.67 mL, 0.50 g, 3.84 mmol) in DCM (9 mL) was added (by syringe pump), over the course of 1 hour, to a solution of triphosgene (352 mg, 1.19 mmol) in DCM (7 mL). The resulting mixture was stirred for 1 h at rt before a mixture of 4-[3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile (1.08 g, 3.58 mmol, see Example G above) and *N*-ethyl di-*iso*-propylamine (0.67 mL, 0.50 g, 3.84 mmol) in DCM (14 mL) was added. Stirring was continued

for a further 20 min, before the solution was concentrated *in vacuo* and the resulting residue purified by flash chromatography, eluting with dichloromethane:methanol (95:5), to give the title compound in 65% yield.

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$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  20.02, 20.16, 29.11, 29.32, 29.46, 31.91, 37.83, 37.89, 48.23, 48.47, 49.36, 49.61, 56.95, 60.26, 60.51, 61.58, 62.077, 65.43, 70.69, 104.06, 115.40, 119.27, 120.77, 133.96, 157.08, 162.21

#### 10 Example 24

7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-(2,2,2-trifluoroethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

The title compound was prepared in 46% yield according to the procedure described in Example 23(b) above, using 2,2,2-trifluoroethylamine in place of 2-aminopropanenitrile.

15

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  29.11, 29.42, 31.79, 42.17, 42.51, 48.36, 49.58, 57.09, 60.45, 61.77, 65.39, 70.76, 104.08, 115.39, 119.23, 123.28, 126.05, 133.93, 157.76, 162.21

20

#### Example 25

7-[3-(4-Cyanophenoxy)-2-hydroxypropyl]-N-[2-oxo-2-(1-piperidiny)-ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

The title compound was prepared in 49% yield according to the procedure described in Example 12(b) above, using piperidine in place of propylamine.

25

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  24.33, 25.41, 26.06, 28.74, 29.29, 29.44, 32.13, 42.67, 43.10, 45.30, 47.99, 48.09, 49.14, 49.28, 57.18, 60.42, 61.90,

65.55, 70.77, 94.22, 103.89, 115.24, 115.43, 119.24, 133.74, 134.02, 158.49, 162.20, 167.42

#### Example 26

5 *N*-(1,3-Benzodioxol-5-yl)-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

The title compound was prepared in 33% yield according to the procedure described in Example 23(b) above, using 1,3-benzodioxol-5-amine in place of 2-aminopropanenitrile.

10

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 162.22, 156.51, 147.47, 143.20, 133.98, 133.83, 119.41, 115.40, 113.68, 107.68, 103.83, 103.59, 100.96, 70.70, 65.98, 61.34, 60.34, 57.87, 49.17, 48.13, 31.52, 29.41, 29.11

15 Example 27

7-[3-(4-Cyanoanilino)propyl]-*N*-[2-oxo-2-(propylamino)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

(a) 4-[(3-Hydroxypropyl)amino]benzonitrile

20 A mixture of 4-fluorobenzonitrile (12.0 g, 99.1 mmol) and 3-amino-1-propanol (59.6 g, 793 mmol) was stirred at 80°C under an inert atmosphere for 3 hours before water (150 mL) was added. The mixture was allowed to cool to rt, and was then extracted with diethyl ether. The organic layer was separated, dried (Na<sub>2</sub>SO<sub>4</sub>), filtered and concentrated *in*  
 25 *vacuo* to yield 17 g (97%) of the title compound as a oil that crystallised upon standing.

(b) 3-(4-Cyanoanilino)propyl 4-methylbenzenesulfonate

A cooled (0°C) solution of 4-[(3-hydroxypropyl)amino]benzonitrile (17 g, 96.5 mmol, from step (a) above) in dry MeCN (195 mL) was treated with triethylamine (9.8 g, 96.5 mmol) and then *p*-toluenesulfonyl chloride (20.2 g, 106 mmol). The mixture was stirred at 0°C for 90 minutes before being concentrated *in vacuo*. Water (200 mL) was added to the residue, and the aqueous solution was extracted with DCM. The organic phase was dried (Na<sub>2</sub>SO<sub>4</sub>), filtered and concentrated *in vacuo*. The resulting residue was purified by crystallisation from *iso*-propanol to yield 24.6 g (77%) of the sub-title compound.

(c) Ethyl 2-{[(7-benzyl-3,7-diazabicyclo[3.3.1]non-3-yl)carbonyl]amino}-acetate

The sub-title compound was prepared in 99% yield according to the procedure described in Example 5(a) above, using 4-[3-(3,7-diazabicyclo[3.3.1]non-3-yl)-2-hydroxypropoxy]benzonitrile (see Example G above) and ethyl 2-isocyanatoacetate in place of 3-benzyl-3,7-diazabicyclo[3.3.1]nonane and *iso*-propyl isocyanate, respectively.

(d) 7-Benzyl-*N*-[2-oxo-2-(propylamino)ethyl]-3,7-diazabicyclo[3.3.1]-nonane-3-carboxamide

The sub-title compound was prepared in 88% yield according to the procedure described in Example 12(b) above, using ethyl 2-{[(7-benzyl-3,7-diazabicyclo[3.3.1]non-3-yl)carbonyl]amino}acetate (from step (c) above) in place of ethyl 2-[(7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]non-3-yl)carbonyl]amino]acetate.



(e) *N*-[2-Oxo-2-(propylamino)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

The title compound was prepared according to the procedure described in Example 5(b) above, using 7-benzyl-*N*-[2-oxo-2-(propylamino)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (from step (d) above) in place of 7-benzyl-*N*-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide.

(f) 7-[3-(4-Cyanoanilino)propyl]-*N*-[2-oxo-2-(propylamino)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of *N*-[2-oxo-2-(propylamino)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (3.35 g, 12.5 mmol, from step (e) above), K<sub>2</sub>CO<sub>3</sub> (6.9 g, 50 mmol) and sodium iodide (0.19 g, 1.25 mmol) in acetonitrile (600 mL) was treated with 3-(4-cyanoanilino)propyl 4-methylbenzenesulfonate (4.2 g, 12.7 mmol, from step (b) above) and stirred under reflux for 5 h, followed by a further 21 h at rt. The mixture was filtered, concentrated *in vacuo* and the crude product so obtained was diluted with water. The aqueous solution was extracted with DCM, which organic layer was separated, dried and concentrated *in vacuo*. The crude product so obtained was purified by chromatography on silica gel, eluting with DCM:MeOH (95:5) to yield 3.08 g (58%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 11.49, 22.85, 25.11, 29.09, 31.03, 40.78, 41.40, 44.80, 48.41, 56.22, 59.32, 97.43, 111.99, 120.97, 133.74, 151.98, 157.92, 170.37

Example 287-{2-[2-(4-Cyanophenoxy)ethoxy]ethyl}-N-ethyl-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide5 (a) 4-[2-(2-Hydroxyethoxy)ethoxy]benzonitrile

A mixture of *p*-cyanophenol (11.9 g, 100 mmol), K<sub>2</sub>CO<sub>3</sub> (15 g, 110 mmol) and chloroethoxyethanol (12.4 g 100 mmol) in CH<sub>3</sub>CN was refluxed for 24 h, then stirred at rt for a further 2 days. The reaction mixture was filtered and concentrated *in vacuo* to give a crude product  
 10 which was purified by chromatography on silica gel (hexane:ethyl acetate (1:1) eluant). This gave 10 g (50%) of the sub-title compound.

(b) 2-[2-(4-Cyanophenoxy)ethoxy]ethyl methanesulfonate

Methanesulfonyl chloride (3.0 g, 26 mmol) was added dropwise to a  
 15 cooled (-5°C) mixture of 4-[2-(2-hydroxyethoxy)ethoxy]benzonitrile (5.0 g, 24 mmol, from step (a) above) and triethylamine (4 mL, 2.9 g, 29 mmol) in DCM (50 mL). After addition was complete, the reaction was allowed to warm to rt over a period of 2 h. The reaction mixture was then washed twice with water, the organic layer separated, dried (Na<sub>2</sub>CO<sub>3</sub>) and  
 20 concentrated *in vacuo* to yield 7 g (100%) of the sub-title compound.

(c) 7-{2-[2-(4-Cyanophenoxy)ethoxy]ethyl}-N-ethyl-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide

A mixture of 2-[2-(4-cyanophenoxy)ethoxy]ethyl methanesulfonate (2 g, 7.0 mmol, from step (b) above), *N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (1.4 g, 7.0 mmol, see Example 6(b) above) and K<sub>2</sub>CO<sub>3</sub> (1.5 g, 10.5 mmol) in MeCN (50 mL) was stirred under reflux overnight. The reaction mixture was then concentrated *in vacuo* and the resulting residue  
 25 purified by flash chromatography on silica gel, eluting with

dichloromethane:methanol (9:1) to yield 0.8 g (30%) of the title compound.

$^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  162.18, 133.92, 119.24, 115.34, 103.92, 69.07,  
5 67.86, 59.52, 58.42, 48.18, 35.68, 30.25, 28.81, 15.69

### Example 29

7-[4-(4-Cyanophenyl)-4-(3,4-dimethoxyphenoxy)butyl]-N-ethyl-3,7-diaza-  
bicyclo[3.3.1]nonane-3-carboxamide

#### (a) 4-[1-(3,4-Dimethoxyphenoxy)-3-butenyl]benzonitrile

A cooled ( $0^\circ\text{C}$ ) mixture of 4-(1-hydroxy-3-butenyl)benzonitrile (14.6 g, 84.3 mmol) and 3,4-dimethoxyphenol (19.5 g, 125.4 mmol) in toluene (500 mL) was treated with tributylphosphine (32.14 mL of 97% purity, 25.6 g, 126.4 mmol), followed by 1,1'-(azodicarbonyl)dipiperidine (31.8 g, 126.4 mmol). After addition was complete, the reaction mixture thickened and the temperature rose to  $15^\circ\text{C}$ . Additional toluene was added (500 mL), and the mixture stirred at rt overnight. The precipitate of tributylphosphine oxide was then removed by filtration and the filtrate concentrated *in vacuo* to give 65.8 g of crude product. This was purified by chromatography on silica gel, eluting with toluene:methanol (98:2) to yield 17.9 g of the sub-title compound.

#### (b) 4-[1-(3,4-Dimethoxyphenoxy)-4-hydroxybutyl]benzonitrile

Borane-methyl sulfide complex (2 M in ether, 11 mL, 22 mmol) was added dropwise to a cooled ( $-5^\circ\text{C}$ ) solution of 4-[1-(3,4-dimethoxyphenoxy)-3-butenyl]benzonitrile (17.6 g, 56.8 mmol, from step (a) above) in dry THF (15 mL) over a period of 15 minutes (during which time the reaction temperature rose to  $0^\circ\text{C}$ ). The resulting mixture was stirred at

between 0 and 10°C for 1.5 h, before being allowed to warm to rt. Stirring was continued for a further 3.5 h at this temperature before water (22 mL) and sodium perborate tetrahydrate (11 g, 66 mmol) were added. The biphasic mixture was stirred for 2 h at rt before the water layer was separated and extracted with ether. The combined organic layers were washed with brine, dried and concentrated *in vacuo*. The resulting residue was purified by chromatography on silica gel, eluting with *iso*-propanol:ethyl acetate:heptane (5:25:70) to yield 14.5 g (77%) of the sub-title compound.

(c) 4-(4-Cyanophenyl)-4-(3,4-dimethoxyphenoxy)butyl methanesulfonate

A solution of methanesulfonyl chloride (3.4 mL, 5.0 g, 44 mmol) in DCM (15 mL) was added slowly to a cooled (-5°C) mixture of 4-[1-(3,4-dimethoxyphenoxy)-4-hydroxybutyl]benzonitrile (11 g, 34 mmol, from step (b) above) and triethylamine (7 mL, 5.2 g, 50.6 mmol) in DCM (50 mL), during which addition the temperature did not rise above 2°C. Stirring was continued at between 0 and 5°C for a further 2 h before water was added. The resulting organic layer was separated, and washed with water, separated again and then dried to give the sub-title compound in 100% yield.

(d) *tert*-Butyl 7-[4-(4-cyanophenyl)-4-(3,4-dimethoxyphenoxy)butyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate

A mixture of 4-(4-cyanophenyl)-4-(3,4-dimethoxyphenoxy)butyl methanesulfonate (522 mg, 1.29 mmol, from step (c) above), *tert*-butyl 3,7-diazabicyclo[3.3.1]nonane-3-carboxylate (307 mg, 1.356 mmol, see Example F above) and K<sub>2</sub>CO<sub>3</sub> (216 mg, 1.56 mmol) in chloroform:acetonitrile (10 mL of 1:1) was stirred at 70°C for 23 h. The reaction mixture was filtered and the filtrate concentrated *in vacuo* to give

708 mg of crude product. This was purified by flash chromatography, eluting with a gradient of toluene:methanol (97:3 to 10:1), to yield 607 mg (88%) of the sub-title compound.

5 (e) 4-[4-(3,7-Diazabicyclo[3.3.1]non-3-yl)-1-(3,4-dimethoxyphenoxy)-butyl]benzonitrile

A cooled (0°C) solution of *tert*-butyl 7-[4-(4-cyanophenyl)-4-(3,4-dimethoxyphenoxy)butyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate (1.92 g, 3.6 mmol, from step (d) above) in ethyl acetate (20 mL) was  
 10 treated with HCl-saturated ethyl acetate (30 mL). The resulting mixture was stirred for 2 h at between 0 and 5°C before being concentrated *in vacuo*. The resulting residue was dissolved in acetonitrile (50 mL) and treated with K<sub>2</sub>CO<sub>3</sub> (3.5 g, 25.2 mmol) and water (2.25 mL). This mixture was stirred for 3h at rt and the solids removed by filtration,  
 15 before the solvent was removed *in vacuo* (with toluene added to effect azeotropic removal of water) to give 1.5 g of the sub-title compound.

(f) 7-[4-(4-Cyanophenyl)-4-(3,4-dimethoxyphenoxy)butyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

20 A solution of 4-[4-(3,7-diazabicyclo[3.3.1]non-3-yl)-1-(3,4-dimethoxyphenoxy)butyl]benzonitrile (109 mg, 0.25 mmol, from step (e) above), in CHCl<sub>3</sub> (1.43 mL) was treated with a solution of ethyl isocyanate (18.6 µL, 16.8 mg, 0.237 mmol) in MeCN (0.5 mL). The resulting mixture was stirred for 30 h. at rt. The solution was then loaded onto an ion-exchange  
 25 solid phase extraction plug (SiO<sub>2</sub>, 0.5 g from ISOLUTE). The plug was washed with CHCl<sub>3</sub> (2.5 mL) and the product then eluted with MeCN (3 x 2.5 mL). This gave the title compound (93 mg, 73%) in a purity better than 90% (as determined by HPLC: UV at 254 nm and ELS detection).

MS (ES)  $m/z$  = 507 ( $M + 1$ )<sup>+</sup>, 505 ( $M - 1$ )<sup>-</sup>

Example 30

5 7-(3-{4-Cyano-2-[(cyclopropylamino)carbonyl]phenoxy}-2-hydroxy-propyl)-*N*-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

(a) 5-Cyano-*N*-cyclopropyl-2-[2-oxiranylmethoxy]benzamide

The sub-title compound was prepared according to the method described  
10 in Example 7(b) above using 2-oxiranylmethyl 3-nitrobenzenesulfonate (prepared analogously to the method described in Example B above).

(b) 7-Benzyl-*N*-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A cooled (0°C) solution of 3-benzyl-3,7-diazabicyclo[3.3.1]nonane (10 g,  
15 46 mmol, see Example E above) in DCM (100 mL) was treated with phenyl isocyanate (4.9 mL, 45 mmol). The mixture was stirred at rt for 30 min. The product formed as white crystals, which were removed by filtration to give 10 g (66%) of the sub-title compound.

20 (c) *N*-Phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A solution of 7-benzyl-*N*-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (10 g, 29.8 mmol, from step (b) above) in ethanol (100 mL) was subjected to hydrogenation, over 10% Pd/C and at ambient pressure, overnight. The catalyst was removed through a pad of Celite® and the  
25 residue was concentrated *in vacuo* to give the sub-title compound in quantitative yield.

(d) 7-(3-{4-Cyano-2-[(cyclopropylamino)carbonyl]phenoxy}-2-hydroxy-propyl)-N-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of 5-cyano-N-cyclopropyl-2-[2-oxiranylmethoxy]benzamide (0.8 g, 3.1 mmol, from step (a) above) and N-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (0.9 g, 3.6 mmol, from step (c) above) in *iso*-propanol:H<sub>2</sub>O (10 mL of 9:1) was refluxed for 180 min. before dichloromethane was added and the solvent removed *in vacuo*. Purification of the resulting residue by flash chromatography, eluting with DCM:MeOH (9:1), gave 1 g (64%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 6.33, 6.56, 23.23, 29.18, 29.51, 31.66, 48.27, 49.60, 53.44, 57.94, 60.51, 65.74, 71.28, 104.93, 113.46, 118.45, 119.54, 119.65, 122.88, 123.27, 128.84, 136.07, 156.44, 159.69, 164.53

Example 31

N-(4-Cyanophenyl)-7-[3-(ethanesulfonyl)propyl]-3,7-diazabicyclo-[3.3.1]nonane-3-carboxamide

(a) 3-(Ethanesulfonyl)propyl 4-methylbenzenesulfonate

Triethylamine (13.36 g, 132 mmol) was added dropwise to a mixture of 3-(ethanesulfonyl)-1-propanol (13.4 g, 88 mmol, Martin-Smith *et al.*, *J. Pharm. Pharmacol.*, **19**, (1967) 649) and *p*-toluenesulfonyl chloride (16.78 g, 88 mmol) in DCM (150 mL), resulting in a mildly exothermic reaction. After addition was complete, the reaction mixture was washed twice with aqueous ammonium chloride solution, the organic layer was then separated, dried, and concentrated *in vacuo*. The resulting residue was recrystallised from diethyl ether/DCM to give 17.9 g (65%) of the sub-title compound.

(b) *tert*-Butyl 7-[(4-cyanoanilino)carbonyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate

A suspension of *tert*-butyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate (2.0 g, 8.8 mmol, see Example F above) in chloroform (15 mL) was treated with 4-isocyanatobenzonitrile (1.53 g, 10.6 mmol). The mixture was stirred at rt for 1.5 h, at which time some solid particles were observed in the mixture. An additional 10 mL of chloroform was added in order to dissolve the particles. Mass spectroscopic analysis of the mixture indicated that the starting materials had been consumed, and so the solvent was removed *in vacuo*. The resulting residue was purified by flash chromatography, eluting with a gradient of DCM:MeCN (5:1 to 2:1) to yield 2.31 g (71%) of the sub-title compound.

(c) *N*-(4-Cyanophenyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A cooled (0°C) solution of *tert*-butyl 7-[(4-cyanoanilino)carbonyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxylate (2.2 g 5.94 mmol, from step (b) above) in ethyl acetate (40 mL) was treated with HCl-saturated ethyl acetate (65 mL) over the course of 30 minutes. The resulting mixture was stirred at rt for a further 4 h before being concentrated *in vacuo* to give 1.8 g (99%) of the hydrochloride salt of the sub-title compound.

(d) *N*-(4-Cyanophenyl)-7-[3-(ethanesulfonyl)propyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of *N*-(4-cyanophenyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (67.6 mg, 0.25 mmol, from step (c) above) and K<sub>2</sub>CO<sub>3</sub> (80 mg, 0.57 mmol) in DMF (0.5 mL) was treated with a solution of 3-(ethanesulfonyl)propyl 4-methylbenzenesulfonate (153 mg, 0.50 mmol, from step (a) above), in MeCN (1.0 mL). The resulting suspension was stirred for 5 days at 50°C before being cooled and filtered. The filtrate



was then added to a ion-exchange solid phase extraction plug (CBA, 2 g from ISOLUTE). After 1 h the plug was washed with  $\text{CHCl}_3$  (3 x 2.5 mL) and the product eluted with  $\text{CHCl}_3$ :MeOH: $\text{Et}_3\text{N}$  (8:1:1) to give the title compound (63.6 mg, 63%) in a purity better than 90% (as determined by HPLC: UV at 254 nm and ELS detection).

MS (ES):  $m/z = 405$  ( $M + 1$ )<sup>+</sup>,  $m/z = 403$  ( $M - 1$ )<sup>-</sup>

### Example 32

10 7-{3-[(2-Cyano-1*H*-indol-4-yl)oxy]-2-hydroxypropyl}-*N*-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of 4-(2-oxiranylmethoxy)-1*H*-indole-2-carbonitrile (1.0 g, 4.7 mmol, Pitha *et al.*, *J. Med. Chem.*, **30** (1987) 612) and *N*-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (1.4 g, 5.5 mmol, see Example 15 30(d) above) in *iso*-propanol: $\text{H}_2\text{O}$  (10 mL of 9:1) was stirred under reflux for 3 h before being concentrated *in vacuo*. The resulting residue was purified by column chromatography on silica gel, eluting with a gradient of DCM:MeOH (99:1 to 97:3), to yield 0.8 g (37%) of the title compound.

20

<sup>13</sup>C NMR ( $\text{CDCl}_3$ ):  $\delta$  29.03, 29.39, 31.27, 48.37, 49.31, 57.89, 60.42, 61.41, 66.07, 70.04, 100.72, 104.39, 105.13, 111.31, 114.95, 117.66, 120.18, 120.30, 123.00, 126.54, 128.84, 138.39, 139.16, 152.55, 156.29

25

Example 337-[(7-Cyano-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide5 (a) 5-Bromo-2-(3-chloro-2-hydroxypropoxy)benzaldehyde

A mixture of 5-bromo-2-hydroxy benzaldehyde (20.1 g, 0.1 mol), epichlorohydrin (25 mL, 0.32 mol) and 6 drops of piperidine was stirred under reflux for 6 h before being concentrated *in vacuo*. The resulting residue was dissolved in chloroform (25 mL) and treated with concentrated HCl (10 mL). The resulting mixture was stirred for 3 h at rt before the organic layer was washed with water, separated, dried and concentrated *in vacuo* to yield 28.2 g (96%) the sub-title compound. This was used directly in the next step without any further purification.

15 (b) 5-Bromo-2-(3-chloro-2-hydroxypropoxy)phenyl formate

A solution of 5-bromo-2-(3-chloro-2-hydroxypropoxy)benzaldehyde (28.2 g, 96 mmol, from step (a) above) in DCM (200 mL) was treated with 3-chloroperoxybenzoic acid (25 g of 70-75% purity, approximately 100 mmol). The resulting exothermic reaction caused the mixture to reflux for 20 min. Stirring was continued for a further 3 days before the mixture was filtered (to remove precipitated 3-chlorobenzoic acid). The filtrate was washed with K<sub>2</sub>CO<sub>3</sub>-solution and water, dried and concentrated *in vacuo* to yield 26.1 g of sub-title compound. This was used directly in the next step without any further purification.

25

(c) (7-Bromo-2,3-dihydro-1,4-benzodioxin-2-yl)methanol

A solution of 5-bromo-2-(3-chloro-2-hydroxypropoxy)phenyl formate (26.1 g, 84 mmol, from step (b) above) in ethanol (100 mL) was treated with a solution of potassium hydroxide (6.1 g of 85% purity,

approximately 92 mmol) in water (10 mL). The resulting mixture was refluxed for 1.5 h before being filtered and concentrated *in vacuo*. The resulting residue was dissolved in ethyl acetate and washed with brine. The organic layer was separated, dried and concentrated *in vacuo* to give 28.8 g of crude product. This was purified by column chromatography on silica gel, eluting with diethyl ether:hexane (70:30), to yield 10.0 g (49.1%) of the sub-title compound.

(d) 3-(Hydroxymethyl)-2,3-dihydro-1,4-benzodioxin-6-carbonitrile

A mixture of (7-bromo-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (10.0 g, 41.2 mmol, from step (c) above) and CuCN (4.0 g, 45.3 mmol) in DMF (10 mL, dried over molecular sieves) was stirred at 170°C for 4.5 h. The reaction mixture was poured into a warm aqueous solution of sodium cyanide (8.10 g, 165 mmol of NaCN in 25 mL H<sub>2</sub>O). The resulting mixture was extracted with toluene and DCM. The combined organic layers were washed with water and then brine, dried and concentrated *in vacuo*. The residue so obtained was crystallised from toluene and DCM to yield 2.8 g (35%) of the sub-title compound.

(e) (7-Cyano-2,3-dihydro-1,4-benzodioxin-2-yl)methyl methanesulfonate

A solution of methanesulfonyl chloride (1.81 g, 15.8 mmol) in dichloromethane (5 mL) was added dropwise to a cooled (0°C) mixture of 3-(hydroxymethyl)-2,3-dihydro-1,4-benzodioxin-6-carbonitrile (2.75 g, 14.4 mmol, from step (d) above) and pyridine (1.26 g, 16 mmol) in DCM (25 mL). After addition was complete, the mixture was stirred at 0°C for 1 h, and then at rt overnight. TLC analysis indicated incomplete reaction after this time, and so further portions of methanesulfonyl chloride (0.4 g, 3.5 mmol) and pyridine (0.5 mL, 0.49 g, 6.2 mmol) were added. The mixture was refluxed for 3.5 h before being washed twice with saturated

Na<sub>2</sub>CO<sub>3</sub> solution, dried and concentrated *in vacuo*. The crude product (4.5 g) so obtained was purified by flash chromatography, eluting with DCM, to give 3.5 g of the sub-title compound, which crystallised on standing.

5

(f) 7-[(7-Cyano-2,3-dihydro-1,4-benzodioxan-2-yl)methyl]-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of (7-cyano-2,3-dihydro-1,4-benzodioxin-2-yl)methyl methane-sulfonate (150 mg, 0.9 mmol, from step (e) above), *N*-ethyl-3,7-diaza-bicyclo[3.3.1]nonane-3-carboxamide (186 mg, 0.94 mmol, see Example 6(b) above), K<sub>2</sub>CO<sub>3</sub> (265 mg, 2.0 mmol) and NaI (14 mg, 0.09 mmol) in CH<sub>3</sub>CN was refluxed for 20 h. The solvent was removed *in vacuo* and the resulting residue treated with DCM and water. The organic layer was separated, dried (Na<sub>2</sub>SO<sub>4</sub>) and concentrated *in vacuo*. The resulting residue was purified by flash chromatography, eluting with DCM:MeOH (95:5) to yield 113.2 mg (34%) of the title compound.

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<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 15.61, 29.19, 30.72, 35.72, 47.78, 58.34, 59.02, 60.64, 67.01, 71.38, 71.49, 71.60, 104.10, 120.76, 120.89, 125.39, 125.79, 143.50, 147.80, 157.46

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Example 34

7-[(2*S*)-6-Cyano-4-(methanesulfonyl)-3,4-dihydro-2*H*-1,4-benzoxazin-2-yl)methyl]-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

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(a) (2*R*)-2-(Hydroxymethyl)-3,4-dihydro-2*H*-1,4-benzoxazine-6-carbonitrile

A mixture of 3-amino-4-hydroxybenzonitrile (25 g, 0.186 mol) and *S*-epichlorohydrin (10.7 g, 0.22 mol) in aqueous ethanol (500 mL of 99%)

was stirred at 60°C for 24 h. The mixture was concentrated *in vacuo* before ethanol (500 mL) was added, followed by K<sub>2</sub>CO<sub>3</sub> (27 g, 0.195 mol). The resulting mixture was refluxed for 1 h before being filtered. The filtrate was concentrated *in vacuo* to give 61 g of a black oil. This was diluted with water (500 mL), and then extracted twice with DCM and ethyl acetate. The combined organic extracts were dried and concentrated *in vacuo* to yield 20 g (57%) of the sub-title compound as yellow crystals.

(b) (2R)-6-Cyano-4-(methanesulfonyl)-3,4-dihydro-2H-1,4-benzoxazin-2-yl)methyl methanesulfonate

Methanesulfonyl chloride (45 g, 0.395 mol) was added dropwise to a cooled (0°C) mixture of (2R)-2-(hydroxymethyl)-3,4-dihydro-2H-1,4-benzoxazine-6-carbonitrile (30 g, 0.158 mol, from step (a) above) and pyridine (200 mL, excess). The mixture was stirred at rt overnight before being concentrated *in vacuo*. The resulting residue was treated with water and crystals of the product were isolated by filtration. These were recrystallised from MeCN to give 29 g of pure material. The mother liquor was concentrated *in vacuo* to give a residue which was crystallised from chloroform to give a further crop (7.5 g) of product. The total yield of the sub-title compound was 36.5 g (67%).

(c) 7-([(2S)-6-Cyano-4-(methanesulfonyl)-3,4-dihydro-2H-1,4-benzoxazin-2-yl)methyl]-N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A solution of (2R)-6-cyano-4-(methanesulfonyl)-3,4-dihydro-2H-1,4-benzoxazin-2-yl)methyl methanesulfonate (1 g, 2.89 mmol, from step (b) above) in MeCN (5 mL) was treated with triethylamine (8 mL, 5.8 g, 57.4 mmol), followed by N-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (0.85 g, 4.33 mmol, see Example 6(b) above). The resulting mixture was stirred at 70°C for 5h, and then at rt overnight. The mixture

was concentrated *in vacuo* and purified by acid/base extraction, followed by flash chromatography, eluting with DCM:MeOH, to yield 100 mg (14%) of the title compound.

<sup>13</sup>C NMR (CDCl<sub>3</sub>): δ 15.63, 28.87, 29.09, 30.48, 35.73, 39.50, 45.96, 47.65, 48.11, 59.03, 59.19, 60.59, 73.40, 104.15, 118.72, 119.90, 124.92, 126.51, 128.92, 150.04, 157.74

### Example 35

7-[2-({2-[4,5-Bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]acetyl}amino)ethyl)-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

#### (a) 4-[(*E*)-1-(4-Cyanobenzoyl)-2-(dimethylamino)ethenyl]benzonitrile

*N,N*-Dimethylformamide dimethylacetal (135.2 g, 0.29 mol) was added dropwise, under an inert atmosphere, to a heated (60°C) solution of 4-[2-(4-cyanophenyl)acetyl]benzonitrile (60.2 g, 0.24 mol, Ashley *et al.*, *J. Chem. Soc.* (1942) **103**, 110) in 1,2-dimethoxyethane. The resulting mixture was then filtered and concentrated *in vacuo* to give a residue that was crystallised from MeOH. This gave 27.9 g (38%) of the sub-title compound.

#### (b) Ethyl 2-[4,5-bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]acetate

A solution of 4-[(*E*)-1-(4-cyanobenzoyl)-2-(dimethylamino)ethenyl]benzonitrile (6.2 g, 20 mmol from step (a) above) in aqueous ethanol (100 mL of 99%) was treated with ethyl 2-hydrazinoacetate hydrochloride (3.5 g, 22.6 mmol). The mixture was stirred at rt overnight before being concentrated *in vacuo*. The resulting residue was diluted with water, which aqueous mixture was extracted with DCM. The organic layer was then separated, dried and concentrated *in vacuo* to give a residue which

was recrystallised from diethyl ether to yield 1.7 g (23.5%) of the sub-title compound.

(c) 2-[4,5-Bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]-*N*-(2-hydroxyethyl)-acetamide

A mixture of ethyl 2-[4,5-bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]acetate (3.9 g, 10.9 mmol, from step (b) above), 2-amino-1-ethanol (1.3 g, 21.8 mmol) and triethylamine (0.8 g, 76 mmol) was stirred at 100°C overnight. Water and DCM were added, the product crystallised and was isolated by filtration to yield 3.53 g of sub-title compound.

(d) 2-[4,5-Bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]-*N*-(2-bromoethyl)-acetamide

A mixture of 2-[4,5-bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]-*N*-(2-hydroxyethyl)acetamide (0.7 g, 1.88 mmol, from step (c) above), *N*-bromosuccinimide (0.75 g, 5.64 mmol) and triphenylphosphine (2.22 g, 8.4 mmol) in DCM (100 mL) was stirred under reflux for 3 h. The reaction mixture was allowed to cool before being washed with water. The organic layer was separated, dried and concentrated *in vacuo* to give a residue that was purified by flash chromatography, eluting with diethyl ether:methanol (95:5), to yield 0.7 g sub-title compound contaminated with triphenylphosphine oxide. This product was used directly in the next step without any further purification.

(e) 7-[2-({2-[4,5-Bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]acetyl}amino)ethyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide

A mixture of 2-[4,5-bis(4-cyanophenyl)-1*H*-pyrazol-1-yl]-*N*-(2-bromoethyl)acetamide (0.7 g, 1.6 mmol, from step (d) above), *N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide (0.32 g, 1.6 mmol, see

Example 6(b) above) and  $K_2CO_3$  (0.55 g, 4 mmol) in acetonitrile (15 mL) was stirred under reflux overnight. Extraction with diethyl ether and water gave an organic layer that was separated, dried and concentrated *in vacuo*. The resulting residue was purified by chromatography on silica gel, eluting with diethyl ether : MeOH (95:5), to yield 0.27 g of the title compound.

$^{13}C$  NMR ( $CDCl_3$ ):  $\delta$  15.77, 29.18, 32.37, 36.13, 48.72, 52.27, 56.32, 109.83, 113.13, 118.27, 118.93, 120.10, 127.80, 131.39, 132.46, 132.73, 134.62, 138.75, 159.14, 167.09

### Example 36

The following compounds (all of which are title compounds of this Example 36) were also prepared, using analogous methods to those described herein:

- 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[(2*R*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[2-(4-cyanophenoxy)ethyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-tetrahydro-2*H*-pyran-2-yl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-(4-cyanophenethyl)-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;



7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-methyl-3,7-diazabicyclo-

- [3.3.1]nonane-3-carboxamide;
- 7-[(2*S*)-3-(4-cyano-2-{[(2-cyanoethyl)amino]carbonyl}phenoxy)-2-hydroxypropyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- methyl (1*R*)-2-(4-cyanophenoxy)-1-({7-[(ethylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl}methyl)ethylcarbamate;
- 7-[(2*R*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-((2*S*)-3-{4-cyano-2-[(methylamino)carbonyl]phenoxy}-2-hydroxypropyl)-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-propionyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[(2*R*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-propionyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[2-(4-cyanophenyl)-2-hydroxyethyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(2-propynyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-(4-cyanophenethyl)-*N*-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- N*-ethyl-7-[(2*S*)-2-hydroxy-3-(4-nitrophenoxy)propyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- methyl (1*S*)-2-(4-cyanophenoxy)-1-({7-[(ethylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl}methyl)ethylcarbamate;
- 7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(cyclopropylmethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- N*-(4-nitrophenyl)-7-(4-oxoheptyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

- 7-[(2*R*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(2-propynyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-(3-{4-cyano-2-[(cyclopropylamino)carbonyl]phenoxy}-2-hydroxypropyl)-*N*-propionyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 5 7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-phenyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[(2*R*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(cyclopropylmethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- tert*-butyl (1*R*)-2-(4-cyanophenoxy)-1-({7-[(propionylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl}methyl)ethylcarbamate;
- 10 7-(3-{4-cyano-2-[(*iso*-propylamino)carbonyl]phenoxy}-2-hydroxypropyl)-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- tert*-butyl 2-(4-cyanophenoxy)-1-({7-[(propionylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl}methyl)ethylcarbamate;
- 15 *tert*-butyl 2-(4-cyanophenoxy)-1-({7-[(*iso*-propylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl}methyl)ethyl(methyl)carbamate;
- 7-[3-(4-cyanophenoxy)-2-(methylamino)propyl]-*N*-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-{3-(4-cyanophenoxy)-2-[methyl(methylsulfonyl)amino]propyl}-*N*-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 20 *N*-(*tert*-butyl)-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[2-amino-3-(4-cyanophenoxy)propyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 25 *tert*-butyl 2-[7-(aminocarbonyl)-3,7-diazabicyclo[3.3.1]non-3-yl]-1-[(4-cyanophenoxy)methyl]ethylcarbamate;
- tert*-butyl 2-(4-cyanophenoxy)-1-({7-[(tetrahydro-2*H*-pyran-2-ylamino)carbonyl]-3,7-diazabicyclo[3.3.1]non-3-yl}methyl)ethylcarbamate;
- N*-(4-cyanophenyl)-7-(4-oxoheptyl)-3,7-diazabicyclo[3.3.1]nonane-3-

- carboxamide;
- 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(2,2-dimethylpropanoyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- N*-(*tert*-butoxy)-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-
- 5 diazabicyclo[3.3.1]nonane-3-carboxamide;
- 2-[7-(aminocarbonyl)-3,7-diazabicyclo[3.3.1]non-3-yl]-1-[(4-cyano-2-methylphenoxy)methyl]ethyl *tert*-butylcarbamate;
- 7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-*iso*-propyl-*N*-methyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 10 *N*-(4-cyanophenethyl)-7-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- N*-(*tert*-butoxy)-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-methyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- N*-(4-cyanophenethyl)-7-(3,4-dimethoxyphenethyl)-3,7-diazabicyclo-
- 15 [3.3.1]nonane-3-carboxamide;
- 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-cyclopropyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[2-amino-3-(4-cyanophenoxy)propyl]-*N*-(*tert*-butyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 20 *N*-[3-(4-cyanophenoxy)propyl]-7-[5-(ethylamino)-5-oxopentyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(3,5-dimethyl-4-isoxazolyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- 7-[3-(4-cyanoanilino)propyl]-*N*-*iso*-propyl-3,7-diazabicyclo[3.3.1]nonane-
- 25 3-carboxamide;
- 7-[4-(4-cyanophenyl)-4-hydroxybutyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;
- ethyl {7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]non-3-yl}carbonylcarbamate;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(4-cyanophenyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-hexyl-3,7-diazabicyclo[3.3.1]-nonane-3-carboxamide;

*N*-(4-butoxyphenyl)-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(3-cyanophenyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

butyl 2-[(7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo-  
[3.3.1]non-3-yl)carbonyl)amino]acetate;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(4-methoxyphenyl)-3,7-diaza-  
20 bicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(3,4-dimethoxyphenethyl)-  
3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(3,4-dimethoxybenzyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

25 7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(3,4,5-trimethoxyphenyl)-3,7-  
diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(3,4-dihydro-2*H*-1,5-benzodioxepin-7-yl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-(2,6-dimethylphenyl)-3,7-

*iso*-propyl {7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo-  
[3.3.1]non-3-yl}carbonylcarbamate;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-{2-[(cyclopropylmethyl)-amino]-2-oxoethyl}-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

10 *N*-(1-cyano-1-methylethyl)-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-  
diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-iso-propyl-3,7-diazabicyclo-  
[3.3.1]nonane-3-carboxamide;

*N*-[cyano(4-fluorophenyl)methyl]-7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[4-(4-cyanophenoxy)-2-hydroxybutyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]-nonane-3-carboxamide;

7-[4-(4-cyanophenyl)butyl]-*N*-propyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[2-amino-4-(4-cyanophenoxy)butyl]-*N*-propyl-3,7-diazabicyclo[3.3.1]-nonane-3-carboxamide;

7-[4-(4-cyanophenyl)butyl]-*N*-ethyl-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

7-[3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-[2-(2-methoxyethoxy)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

5 *N*-(4-cyanophenyl)-7-(3,3-dimethyl-2-oxobutyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

*N*-(4-cyanophenyl)-7-(3,4-dimethoxyphenethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

10 *N*-(4-cyanophenyl)-7-(cyclopropylmethyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

*N*-(4-cyanophenyl)-7-[2-(2-methoxyethoxy)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

*N*-(4-cyanophenyl)-7-[2-(2,3-dihydro-1,4-benzodioxin-6-yl)-2-oxoethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide;

15 7-[3-(4-acetyl-1-piperazinyl)propyl]-*N*-(4-cyanophenyl)-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide; and

7-[(2*S*)-3-(4-cyanophenoxy)-2-hydroxypropyl]-*N*-[2-oxo-2-(propylamino)ethyl]-3,7-diazabicyclo[3.3.1]nonane-3-carboxamide.

## 20 Example 37

Title compounds of the above Examples were tested in Test A above and were found to exhibit  $D_{10}$  values of more than 6.0.

## Abbreviations

25

AcOH = acetic acid

ADDP = 1,1'-(azodicarbonyl)dipiperidine

aq. = aqueous

atm. = atmospheres

	CBz =	benzyloxycarbonyl
	CDI =	carbonyl diimidazole
	Bu =	butyl
	DCM =	dichloromethane
5	DMF =	dimethylformamide
	DMSO =	dimethylsulfoxide
	Et =	ethyl
	EtOAc =	ethyl acetate
	EtOH =	ethanol
10	ESI =	electron spray interface
	eq. =	equivalents
	FAB =	fast atom bombardment
	h =	hours
	IPA =	<i>iso</i> -propanol
15	<i>i</i> -PrOH =	<i>iso</i> -propanol
	LC =	liquid chromatography
	HPLC =	high performance liquid chromatography
	<i>m</i> CPBA =	<i>meta</i> -chloroperbenzoic acid
	Me =	methyl
20	MeCN =	acetonitrile
	MeOH =	methanol
	mesyl =	methanesulfonate
	min. =	minutes
	Ms =	mesylate
25	MS =	mass spectroscopy
	NADPH =	nicotinamide adenine dinucleotide phosphate, reduced form
	NMR =	nuclear magnetic resonance
	OSu =	O-succinyl



	Pd/C =	palladium on carbon
	<i>p</i> TSA =	<i>para</i> -toluenesulfonic acid
	rt. =	room temperature
	satd. =	saturated
5	TEA =	triethylamine
	THF =	tetrahydrofuran
	tlc =	thin layer chromatography
	TMS =	tetramethylsilane

10

Prefixes *n*-, *s*-, *i*-, *iso*-, *t*- and *tert*- have their usual meanings: normal, *iso*, secondary and tertiary.